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PROCESS FEASIBILITY STUDY IN SUPPORT OF SILICON MATERIAL TASK I

JPL Contract No. 954343

Silicon Material Task Low-Cost Solar Array Project to JET PROPULSION LABORATORY CALIFORNIA INSTITUTE OF TECHNOLOGY

by Carl L. Yaws Ku-Yen Li Jack R. Hopper C. S. Fang Keith C. Hansen

February 6, 1981

This work was performed for the Jet Propulsion Laboratory, California Institute of Technology, under NASA Contract NAS7100 for the U.S. Department of Energy, Division of Solar Energy.

The JPL Low-Cost Solar Array Project is funded by DOE and forms part of the DOE Photovoltaic Conversion Program to initiate a major effort toward the development of low-cost solar arrays.

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ABSTRACT

The Low-Cost Solar Array (LSA) Project at Jet Propulsion Laboratory (JPL) in Pasadena, California is being funded by the Department of Energy (DOE) for effective cost reduction in the production of silicon for solar cells. This study reports work performed at Lamar University in support of the LSA Project and presents results for process system properties, chemical engineering and economic analyses of the new technologies and processes being developed for the production of lower cost silicon for solar cells.

Analyses of process system properties are important for chemical materials involved in the several processes under consideration for semiconductor and solar cell grade silicon production. Major physical, thermodynamic and transport property data are reported for the following silicon source and processing chemical materials

- · Silane
- · Silicon Tetrachloride
- · Trichlorosilane
- · Dichlorosilane
- · Silicon Tetrafluoride
- · Silicon

The property data are reported for critical temperature, critical pressure, critical volume, vapor pressure, heat of vaporization, heat capacity, density, surface tension, viscosity, thermal conductivity, heat of formation and Gibb's free energy of formation. The reported property data are presented as a function of temperature to permit rapid usage in research, development and production engineering.

Chemical engineering analyses involving the preliminary process design of a plant (1000MT/yr capacity) to produce silicon via the technology under consideration was accomplished for the following processes:

- · UCC Silane Process for Silicon
- · BCL Process for Silicon Case A
- · BCL Process for Silicon Case B
- · Conventional Polysilicon Proc: (Siemens Technology)
- SiI_A Decomposition Process
- DCS Process (Dichlorosilane)

Major activities in the chemical engineering analyses included base case conditions, reaction chemistry, process flowsheet, material balance, energy balance, property data, equipment design, major equipment list, production labor and forward for economic analysis. The process design package provided detailed

data for raw materials, utilities, major process equipment and production labor requirements necessary for polysilicon production in each process.

Using detailed data from the process design package, economic analyses for a 1000MT/yr silicon plant were accomplished for the processes under consideration for production of lower cost silicon. Primary results issuing from the economic analyses included plant capital investment and product cost which are useful in identification of those processes showing promise for meeting project cost goals.

Cost and profitability results issuing from the chemical engineering and economic analyses are summarized below:

Process	Product Cost, \$/kg (1980 dollars)	Sales Price, \$/kg (1980 dollars)
•UCC Silane Process for Silicon	9.88	13.00 @ 15% DCF
*BCL Process for Silicon - Case A	12.08	13.28 @ 5% DCF
•BCL Process for Silicon - Case B	11.07	13.14 @ 10% DCF
·Conventional Polysi- licon Process Siemens Technology)	53.77	La - 07 fe La
•SiI _A Decomposition Process	62.50	71.48 @ 5% DCF

For the summary tabulation, the product cost represents all cost associated with producing silicon including direct manufacturing cost, indirect manufacturing cost, plant overhead and general expenses. The sales price includes a profit for he company measured in terms of DCF (discounted cash flow) rate of return on the capital investment that the company spent in going into the business.

The cost and profitability analysis results of \$9.66 and \$13 per kg (1980 dollars) at 15% DCF for producing silicon by the UCC silane process (Union Carbide Corporation) indicate that this new technology for producing polysilicon shows good promise for meeting the cost goal of \$14 per kg of silicon naterial (1980 dollars) for solar cells.

For the BCL process - Cases A and B (Battelle Columbus Laboratories), cost and profitability results are \$11.07 - 12.08 and \$13.14 - 13.28 per kg (1980 dollars) at 5 - 10% DCF rate of return. These results indicate that this new technology for producing polysilicon shows promise for meeting the cost goal of \$14 per kg of silicon material (1980 dollars) for solar cells.

For the conventional polysiticon process, the cost analysis is based on a poly plant constructed in the 1980's (1985 or earlier) since several existing plants producing semiconductor grade polysilicon in the United States were constructed in the 1980's. The operating costs for the plant are applicable to the time period of interest such as 1980. The average product cost, \$53.77 per kg (1980 dollars), for the conventional polysilicon process corresponds to intermediate electrical costs (3.15 c/kw hr for 1980). These costs results for the conventional polysilicon process indicate that this Siemens technology using trichlorosilane for producing polysilicon does not show promise for meeting the cost goal of \$14 per kg of silicon material (1980 dollars) for solar cells.

The cost and profitability results of \$62.5 and \$71.48 per kg (1980 dollars) at 5% DCF rate of return for the SiI₄ decomposition process indicate that this new technology for producing polysilicon does not show promise for meeting the cost goal of \$14 per kg of silicon material (1980 dollars) for solar cells.

Using a hot wire method, gas phase thermal conductivity values of argon, hydrogen, silane, dichlorosilane, trichlorosilane, tetrachlorosilane and tetrafluorosilane were experimentally desermined between 25°C and 350°C. Comparison of the values obtained in the study for argon and hydrogen with previously reported values indicated that the values should be accurate to 12° throughout the temperature rap, e.

Using a transpiration technique, gaseous viscosity values for nitrogen, dichlorosilane, trichlorosilane, and tetrafluorosilane were experimentally determined between 40°C and 200°C. Comparison of the values obtained in the study for nitrogen with previously reported viscosity values indicate that the values obtained are accurate to +2° throughout the temperature range

Studies were conducted to develop a method of generating silicon tetrafluoride from hexafluorasilicic acid, a readily available by-product of the phosphate fertilizer industry. Conditions for the efficient precipitation of two SiF precursors (Na₂SiF₆ and BaSiF₆) were determined. These precursors were then thermally decomposed to generate SiF₄. Parameters such as temperature, heating time, and flow rate necessary for efficient production of SiF₄ were determined.

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INTRODUCTION - BACKGROUND

The Low-Cost Solar Array Project (LSA) of Jet Propulsion Laboratory (JPL) had its beginning in 1975, and was concerned with achieving our national solar energy goal (1) to "develop at the earliest feasible time those applications of solar energy that can be made economically attractive and environmentally acceptable as alternate energy sources." Solar cell grade silicon for photovoltaic systems will need to be produced in great volume at considerable reduced prices to accomplish this significant energy goal.

The Low-Cost Solar Array (LSA) Project at Jet Propulsion Laboratory (JPL) in Pasadena, California is being funded by the Department of Energy (DOE) for effective cost reduction in the production of silicon for solar cells. An important overall objective of the project is to reduce the cost of electricity produced with solar cells from today's \$10-25 per W (Peak) to \$0.70 per W (1980 dollars) by 1986. Cost reductions for solar cells are allocated to major tasks encompassing everything from initial silicon praction to final array assembly. The cost goal for the silicon material that goes into solar cells is about \$14 per kg of material (1980 dollars).

Semiconductor grade silicon which is currently produced via the conventional Siemens process by several major manufacturers (Dow-Corning, Monsanto, Motorola, Texas Instruments and Great Western) in the United States is too expensive to meet the silicon material cost goal. Lower cost silicon is needed for solar cells. Alternate processes that depart from the conventional process need to be developed by several concerns to produce a less costly silicon material.

Process evaluation - which is a very useful tool in research and development - is useful in investigation of such alternate processes for solar cell grade silicon. The planning and implementation of a research and development program involves decision making on what work can be left out with least jeopardy to short and long term consequences and what work should be pursued with the best chance for success in achieving short and long term goals. Early process evaluation investigation including preliminary economic evaluation aids the decision making involved in whether to commit extra funds to carryout a project from research to large scale plant.

^{1.} ERDA, National Solar Energy Program, Industrial Briefing, NASA/JPL Low-Cost Silicon Solar Array Project, NASA Headquarters, Washington, D.C. (February 5, 1975).

The early study particularly minimizes the risks involved in the process development from early research to large scale plant. The process evaluation investigation should be initiated with the very inception of the research project and continued throughout its life until the project is proved successful or abandoned because it cannot effectively meet the financial and product purity goals.

In research and development, a screening out is required for those projects and processes which are believed to be unsound or least attractive. Economics dictate that the money should not be wasted on projects which may turn out to be useless. The many alternate projects and processes which are available necessitate the effective use of a screening procedure, not to locate a foolproof venture, but to try to select the best possible project.

Process evaluation investigation may effectively deal with a complete process or part of a process. Major cost areas of a process and profitability potential of a proposed process may be pinpointed. It is also equally valuable in comparing alternate processes and in the selection of processes with the best technical and economic features.

A typical sequence for process selection is presented in Figure 1-1. The process evaluation activities are shown in relation to their usefulness in the selection of a process for scaleup to pilot plant and large scale plant. These process evaluation activities (system properties, chemical engineering and economic analysis tasks) may be effectively utilized in the investigation of alternate processes for low cost, high volume production of silicon suitable for solar cells.

In this process feasibility study in support of Silicon Material Task I of the LSA, the proposed scope of work is to perform investigations and analyses of processes for the low cost, high volume production of silicon suitable for solar cells. The objective of this program is to validate the commercial practicality of these alternate processes based on the following process evaluation criteria:

- Analyses of Process-System Properties Chemical Engineering Analyses
- Economic Analyses

Each of these evaluation criteria is focused on the production of solar cell grad, silicon at greatly reduced cost.

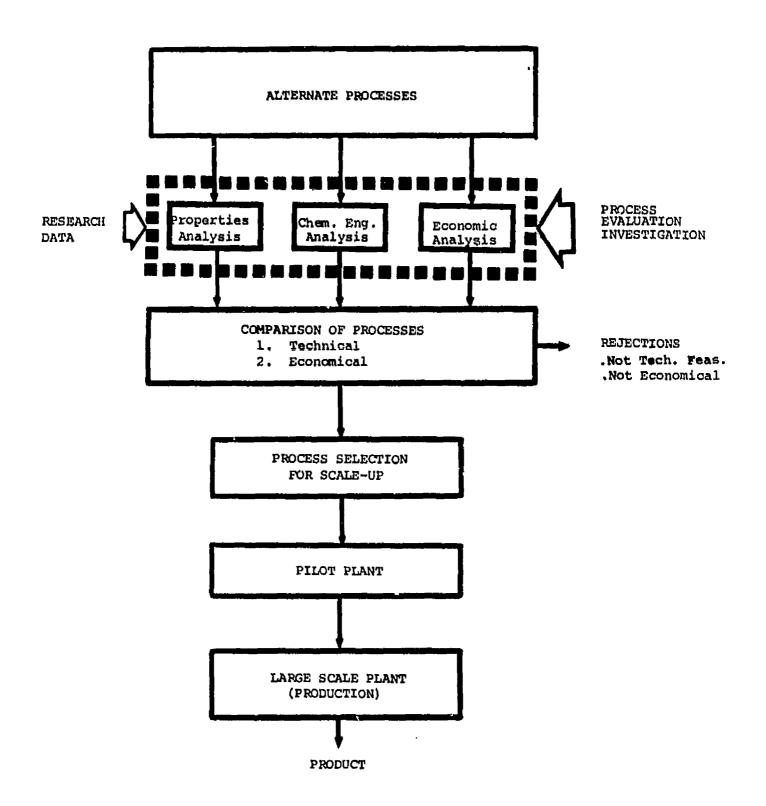


Figure 1-1 TYPICAL SEQUENCE FOR PROCESS SELECTION

PROCESS SYSTEM PROPERTIES ANALYSES

2.1 Silane Properties

Critical Properties (Table 2.1-1)

Experimental data for the critical temperature and pressure of silane are available (2, 15, 20, 22, 25, 27, 29, 45, 47, 49, 50, 51). However, all sources cite Adwentowski (51), who prepared his sample from MgoSi and used the fraction boiling at -116°C. the boiling point of silane is generally accepted as -112°C, these data may not be completely reliable. The critical compressibility factor, $Z_{\rm c}$, was esti-

mated by the Garcia-Barcena method:

$$Z_c = f(T_b) - g(T_b/M)$$
 (2.1-1)

where T_b = boiling point, °K; and M = molecular weight g/g - mol. The terms f (T_b) and g (T_b/M) are shown as a nomograph (29). Reid and Sherwood tested this equation using sixteen inorganic compounds and found an average deviation of 1.8%. The accuracy of the correlation was tested by applying it to similar inorganic and organic compounds (NH₃, N₂H₄, B₂H₆, CH₁). Average deviation was 3.6% for the compounds tested.

The critical volume was found by the real gas relation:

$$V_c = Z_c RT_c / P_c \qquad (2.1-2)$$

using the Adwentowski data and the estimated value of Z.

Vapor Pressure (Figure 2.1-1)

Observed vapor pressure data for silane are available (2, 13, 15, 18, 20, 25, 27, 36, 45, 51, 60) over nearly the entire liquid phase from melting point (mp) to boiling point (bp) to critical point (cp). The available data were correlated with the least squares technique for vapor pressure as a function of temperature using the following correlation relation (64):

$$\log P_v = A + \frac{B}{T} + C \log T + DT + ET^2$$
 (2.1-3)

Average deviations were less than 3.5%. Greater deviations were encountered with other vapor pressure equations. For example, average absolute deviations exceeded 38% for the Cox-Antoine type equation.

Heat of Vaporization (Figure 2.1-2)

Heat of vaporization data for silane are available only at the boiling point (2, 21, 22, 23, 27, 41).

These data vary less than 1%. Watson's correlation (27, 29) was used to extend the heat of vaporization over the entire liquid phase:

over the entire liquid phase: $\Delta H_{v} = \Delta H_{v1} \left(\frac{T_{c} - T}{T_{c} - T_{1}} \right)^{n} \qquad (2.1-4)$

where ΔH_{v_1} is the heat of vaporization at the boiling point (T_1^v) and n = 0.38.

Heat Capacity (Figures 2.1-3 and 2.1-4)

Heat capacity of the ideal gas at atmospheric pressure is primarily based on structural and spectral data. Values from the various sources (5, 16, 20, 22, 25, 39, 44, 52) are in excellent agreement with differences less than 1%.

Liquid heat capacity data (6) are available in the mp-bp temperature interval. The data were extended to cover the full liquid phase with the density relation: liquid heat capacity × density = constant. The constant value was .2895. Testing of the relationship with the available data produced average deviation of 7%.

Density (Figure 2.1-5)

Liquid density data for silane are available (2, 15, 18, 23, 25, 35, 48, 52) from the melting point to the boiling point. The Yaws-Shah equation (62) for density of the saturated liquid was used to extend the data to the critical point:

$$\rho = AB^{-(1-T_r)^{2/7}}$$
 (2.1-5)

where ρ = density, g/cm^3 , T_r = reduced temperature, T/T_c , A, B = correlation parameters. The correlation parameter values for silane are A = 0.2447 and B = 0.3137. Average deviation of calculated and experimental data was 1.48%.

Surface Tension (Figure 2.1-6,

Data for surface tension (7) are available from the melting point to the boiling point. These data were extended using the Othmer relation (29):

extended using the Othmer relation (29):

$$\sigma = \sigma_1 \left(\frac{T_c - T}{T_c - T_1} \right) \qquad (2.1-6)$$

where σ_1 = surface tension at T_1 , dynes/cm; T_1 = critical emperature, °K; T_1 = temperature, °K; and T_1 = the correlation parameter, 1.2. Deviations between data and correlated values were less than 1%.

Viscosity (Figures 2.1-7 and 2.1-8)

The Stiel and Thodos correlation (29) was used to

augment limited data on gas viscosity (2, 15, 20, 23, 25, 52, 53, 57) at atmospheric pressure. All data sources cite Rankine (57) who made his measurement in 1922 at 15° and 100°C. Deviations between data and correlation were less than 1% for the two data values.

Liquid viscosity data are available (30) in the temperature range between the melting point and boiling point. The data were extended to cover the entire liquid range with the following correlation (63) for viscosity of the saturated liquid as a function of temperature:

$$\log \mu_{L} = A + B/T + CT + DT^{2}$$
 (2.1-7)

Correlation values and data were in good agreement with average absolute deviation of 1.4%.

Thermal Conductivity (Figures 2.1-9 and 2.1-10)

Gas thermal conductivity for silane was estimated by the modified Eucken correlation for polyatomic gases. The Eucken correlation agrees well with Svehla (40); deviations were less than 1%. There are no experimental data available for gas thermal conductivity.

Liquid thermal conductivity for silane was estimated with the modified Stiel and Thodos relation (29). The correlation was tested with experimental data for methane. The average deviations were less than 17%. The deviations for silane are probably in the same range. The presented results are intended to represent correct order-of-magnitude values.

Heat of Formation and Free Energy of Formation (Figures 2.1-11 and 2.1-12)

Values for the heat of formation, ΔH_f , and free energy of formation, ΔG_f , are available from American (39) and Russian (12) sources. Average values were selected. The deviation between data and selected results was 0.2 K cal/mol.

TABLE 2.1-1--CRITICAL CONSTANTS AND PHYSICAL PROPERTIES OF SILANE

<u>Identification</u>	Silane
Formula	SiH ₄
State (std. cond.)	gas (colorless)
Molecular Weight, M	32.12
Boiling Point, Th, •C	-111.9
Melting Point, Tm, •C	-184.7
Critical Temp, T., •C	-3.5 (Questionable Value)
Critical Pressure, Pc, atm	47.8 (Questionable Value)
Critical Volume, V _c , cm ³ /gr mol	130.06 (Estimated)
Critical Compressibility Factor, Z _c	0.281 (Estimated)
Critical Density, pc gr/cm	0.247 (Estimated)
Acentric Factor (Ω)	0.0774

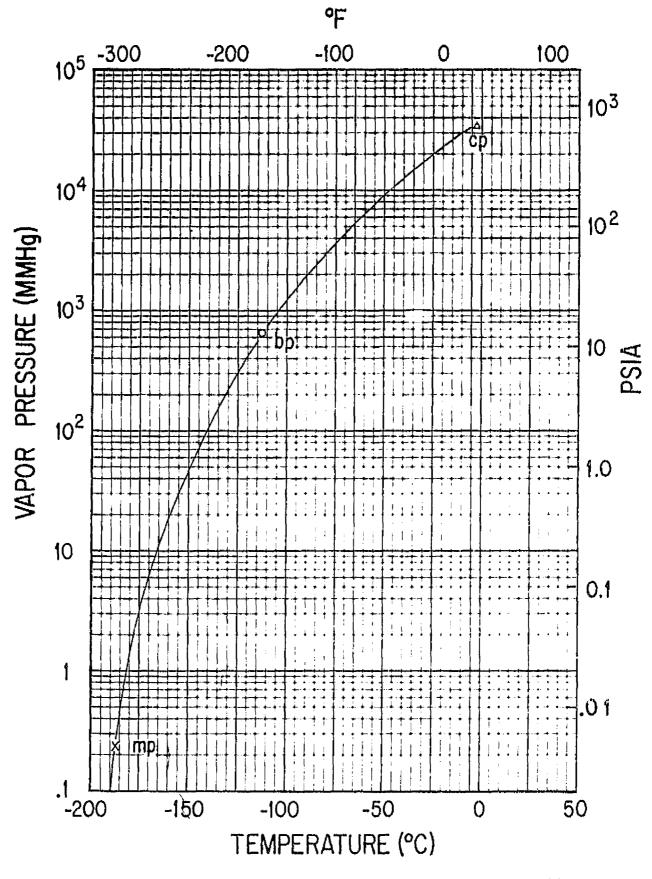


Figure 2.1-1 Vapor Pressure vs. Temperature for Silane

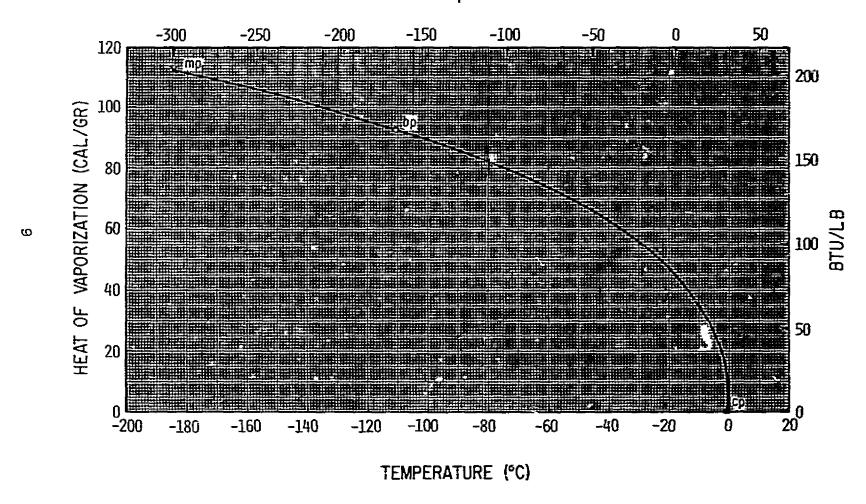


Figure 2.1-2 Heat of Vaporization vs. Temperature for Silane

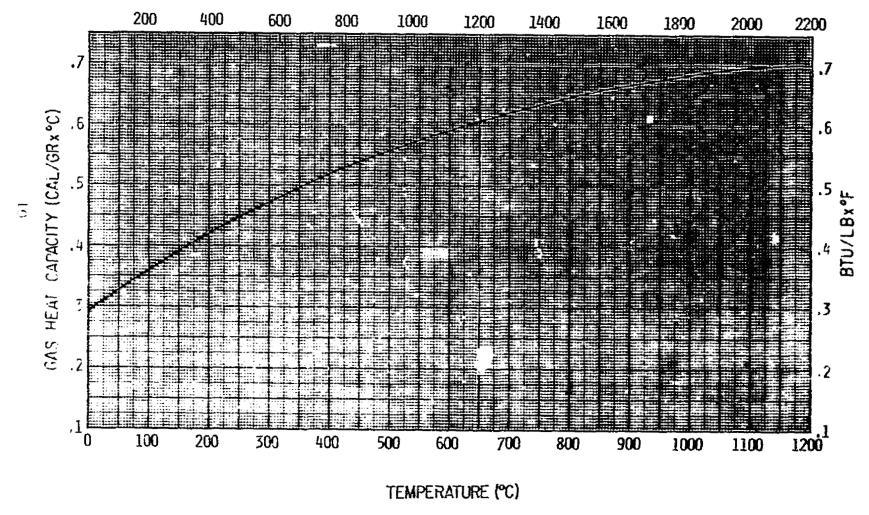


Figure 2.1-3 Gas Heat Capacity vs. Temperature for Silane

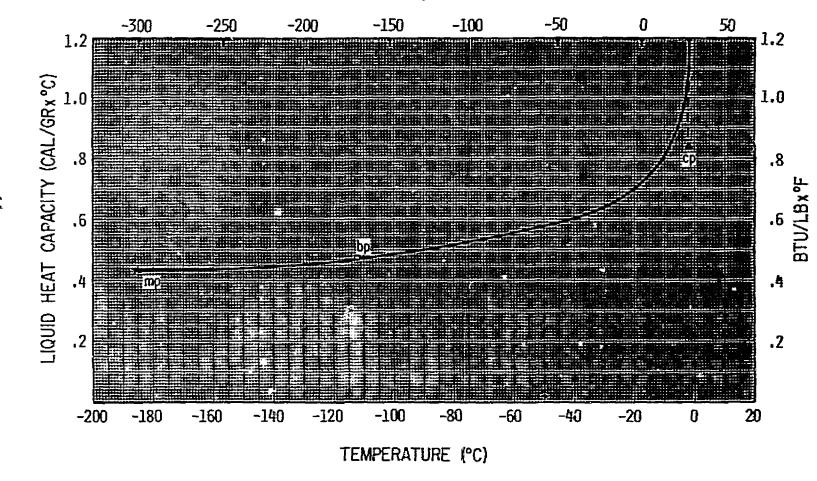


Figure 2.1-4 Liquid Heat Capacity vs. Temperature for Silane

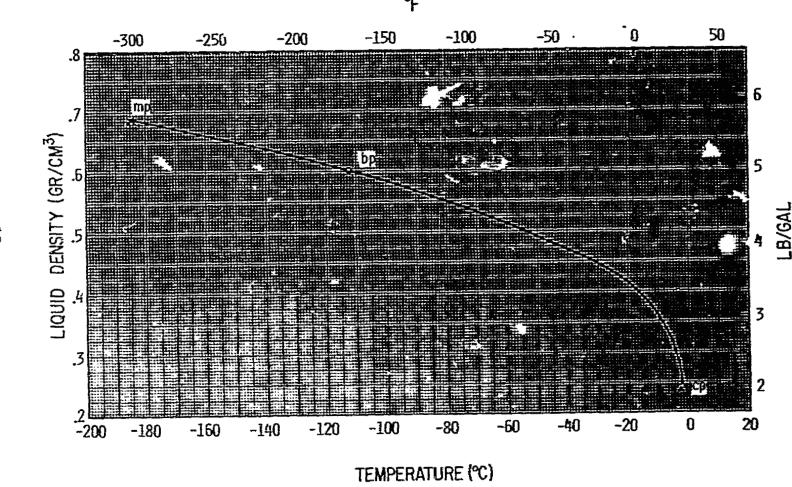


Figure 2.1-5 Liquid Density vs. Temperature for Silane

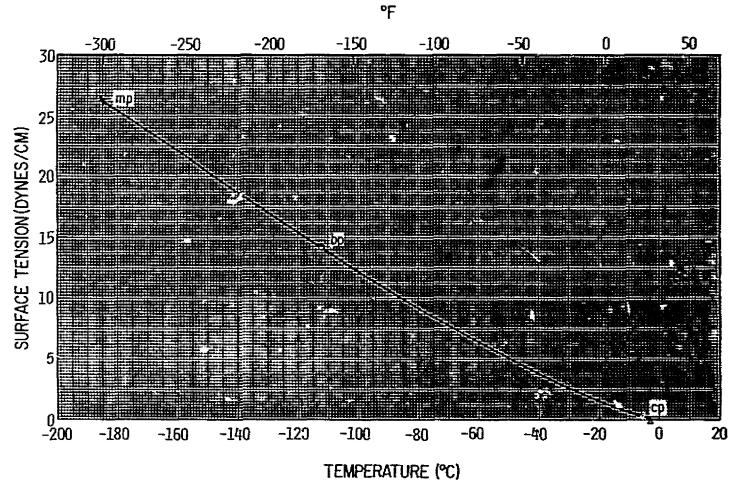


Figure 2.1-6 Surface Tension vs. Temperature for Silane

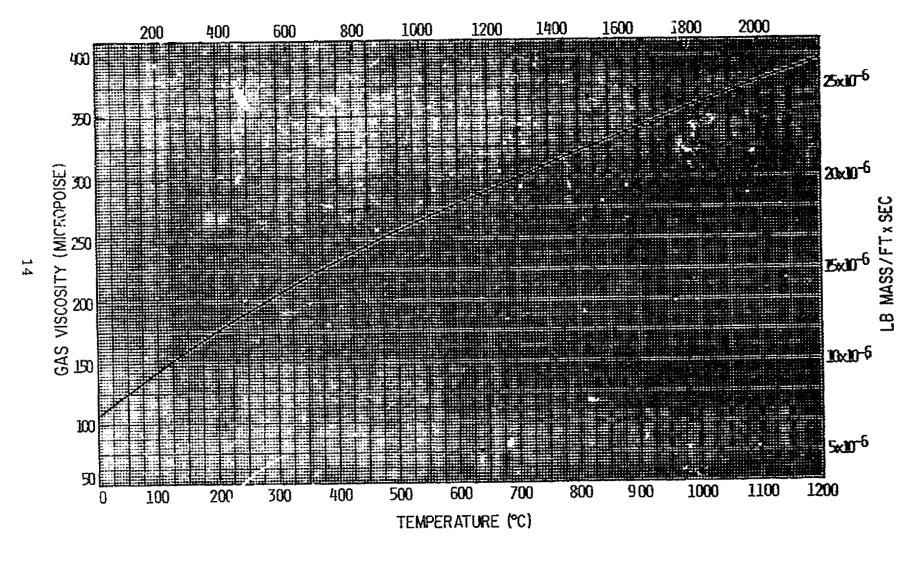


Figure 2.1-7 Gas Viscosity vs. Temperature for Silane

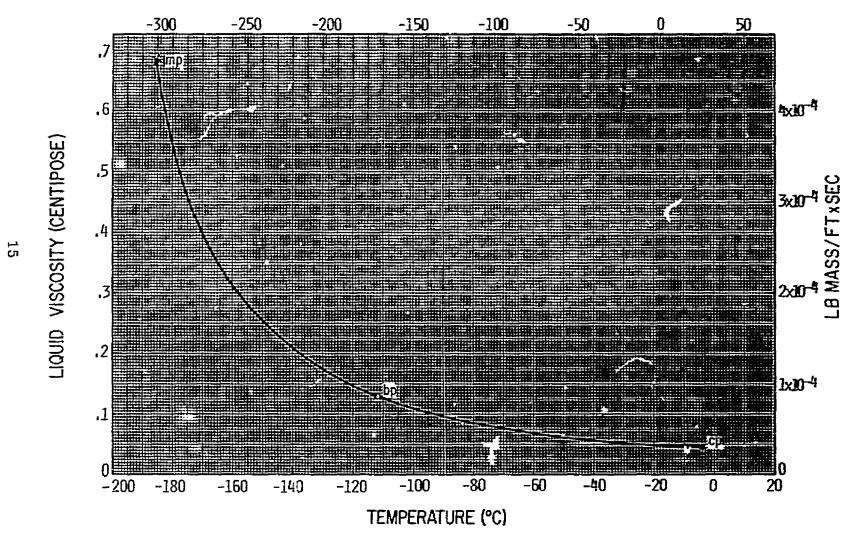


Figure 2.1-8 Liquid Viscosity vs. Temperature for Silane

Figure 2.1-9 Gas Thermal Conductivity vs. Temperature for Silane

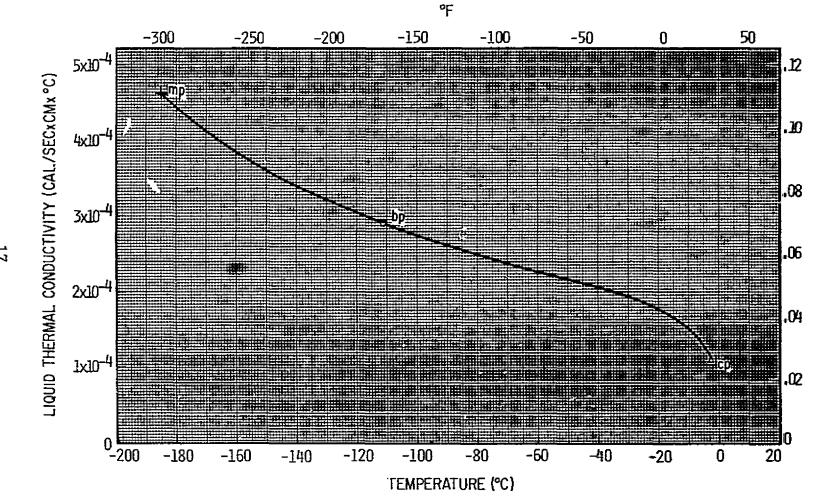


Figure 2.1-10 Liquid Thermal Conductivity vs. Temperature for Silane

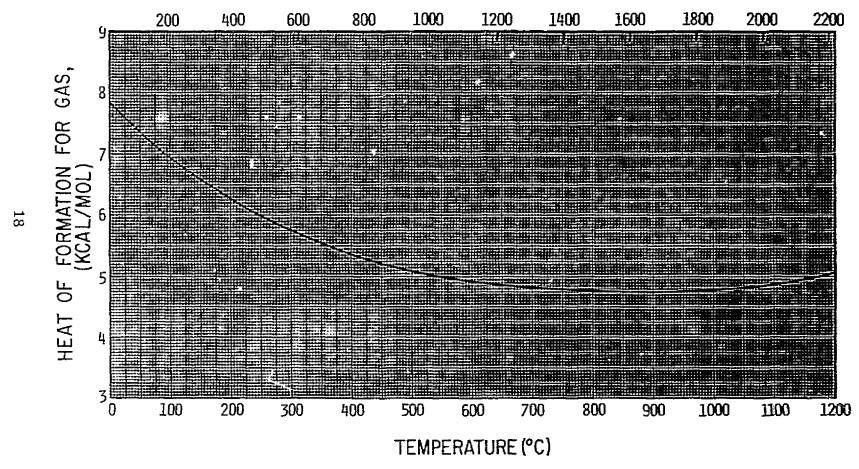


Figure 2.1-11 Heat of Formation vs. Temperature for Silane

°F

Figure 2.1-12 Free Energy of Formation vs. Temperature for Silane

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2.2 Silicon Tetrachloride Properties

Critical Properties (Table 2.2~1)

Experimental results for the critical temperature, pressure and volume of silicon tetrachloride are available (B5, B8, B9, B11, B32, B33, B35, B36, B44, B50, B56, B59, B82, B83). The results among the several investigators are in general agreement. Deviations from the selected values are 1.71%, 0.5%, and 10.8% respectively for critical temperature, pressure and volume.

The critical compressibility factor, $\mathbf{Z}_{\mathbf{C}}$, was calculated using the following equation:

$$Z_{C} = P_{C}V_{C}/RT_{C} \qquad (2.2-1)$$

Also given in the table are values for the acentric factor, $\boldsymbol{\omega}$ which is defined by:

$$\omega = -\log P_r - 1.000 \text{ (at } T_r = 0.70)$$
 (2.2-2)

The acentric factor is an important parameter in generalized thermodynamic correlations involving virial coefficients, compressibility factor, enthalpy and fugacity.

Vapor Pressure (Figure 2.2-1)

Experimental vapor pressure data for silicon tetrachloride are available (B7, B22, B24, B27, B30, B32, B43, B53, B78, B103) from slightly above the melting point (mp) to boiling point (bp) and at the critical point (cp). Available data were extrapolated using the YSSP vapor pressure correlation (B102):

$$\log P_V = A + \frac{B}{T} + C \log T + DT + ET^2$$
 (2.2-3)

where

P, " vapor pressure of saturated liquid, mm of Hg

A, B, C, D, E = correlation constants for chemical compound

The correlation constants (A, B, C, D and E) were determined using a generalized least squares computer program for minimizing deviation of

calculated and experimental data values screened from the literature. Average absolute deviation was about 0.7% for the fifty-eight data points.

Heat of Vaporisation (Figure 2.2-2)

Heat of vaporization data for silicon tetrachloride are available only at the boiling point (B5, B11, B22, B30, B36, B65, B82, B86). Watson's correlation was used to extend the heat of vaporization over the entire liquid phase:

$$\Delta H_{V} = \Delta H_{V_{1}} \left[\frac{T_{C} - T}{T_{C} - T_{1}} \right]^{n}$$
 (2.2-4)

where $\Delta H_{\mathcal{Y}_1}$ is the heat of vaporization at the boiling point (T₁) and n = 0.38.

Heat Capacity (Figures 2.2-3 and 2.2-4)

Heat capacity data for silicon tetrachloride as ideal gas at low pressure are available (B3, B10, B17, B20, B28, B32, B34, B45, B52, B67, B73, B76, B82, B84, B86, B91). The values, which are primarily based on structural and spectral measurements, are in close agreement.

The heat capacity data for the gas phase were correlated by a series expansion in temperature

$$C_{D} = A + BT + CT^{2} + DT^{3}$$
 (2.2-5)

where $C_{\rm p}$ - heat capacity of ideal gas at low pressure, cal/(g-mol)(°K); A, B, C and D = characteristic constants for the chemical compounds; and T = temperature, °K. Average absolute deviation is about 0.6%.

Liquid heat capacity data are available (B5, B22, B28, B30, B26, B43, B52, B60, B65, B76, B77, B82, B104) in the mp-bp temperature interval. The data were extended to cover the entire liquid phase with the relation:

liquid heat capacity x density = constant
$$(2.2-6)$$

The constant value was 0.3054. Testing of the relationship with the available data produced average deviation of 4%.

Density (Figure 2.2-5)

Liquid density data are available (B1, B49, B50, B59, B65, B77, B79, B81, B82) from near the melting point to the critical point. The Yaws-Shah equation (B100, B107) for denisty of the saturated liquid was used to extend the data to the critical point:

$$\rho = AB^{-(1-T_r)^{2/7}}$$
 (2.2-7)

The agreement of calculated and experimental values was very good with average absolute deviation of only 0.44%.

Surface Tension (Figure 2.2-6)

Data for surface tension (B5, B22, B27, B49, B82) are available in the melting point to boiling point temperature range. The data were extended using the Othmer realtion:

$$\sigma = \sigma_1 \left[\frac{T_{\sigma} - T}{T_{\sigma} - T_1} \right]^n \tag{2.2-8}$$

where $\sigma_1 = surface tension at T_1, dynas/cm$

T = critical temperature, °K

T = temperature, *K

and n = the correlation parameter, 1.14. Testing of the relationship with the available data produced average deviation of less than 1%.

Viscosity (Figure 2.2-7 and 2.2-8)

The modified Yoon and Thodos correlation (B105, B106) was used to augment limited data (B36, B51) on gas viscosity at low pressure:

$$\eta \xi = 4.610 \text{ T}_{\perp}^{0.618} - 2.04 \text{ e}^{-0.449 \text{ T}} + 1.94 \text{ e}^{-4.058 \text{ T}} + 0.1 (2.2-9)$$

where
$$\xi = T_c^{1/6} M^{-1/2} P_c^{-2/3}$$

The deviation between data and correlation was 2%.

Liquid viscosity data are available (B5, B8, B32, B36, B49, B51, B107) in both mp-bp and bp-cp temperature ranges. The data were extended to cover the entire liquid range with the following correlation (B107) for viscosity of the saturated liquid as a function of temperature.

$$\log \mu_{T} = A + B/T + CT + DT^{2}$$
 (2.2-]0)

Average deviation between correlation and data was less than 3%.

Thermal Conductivity (Figure 2.2-9 and 2.2-]0)

Gas-phase thermal conductivity data are available (B75, B109) in the temperature range of 70 to 300°C. The data were correlated by a series expansion in temperature (B107):

$$\lambda_{G} = A + BT + CT^{2} + DT^{3}$$
 (2.2-1)

where λ_G = gas thermal conductivity, cal/cmxsecx°C; A, B, C and D = characteristic constants for the chemical compounds and T = temperature, °K. The extimates of Svehla (B73) agree with the above correlation. The deviation between data and correlation values was 1.10%.

Thermal conductivity for liquid phase is available (B25) at only one temperature (32°C). The modified Stiel and Thodos equation (B107) was used to cover the entire saturated-liquid phase.

Heat and Free Energy of Formation (Figures 2.2-11 and 2.2-12)

Values for the heat of formation (ΔH_f) and Gibbs' free energy of formation (ΔG_f) for the ideal gas are available from American (B72) and Russian (B17, B64) sources. American values were selected.

TABLE 2.2-1

CRITICAL CONSTANTS AND PHYSICAL PROPERTIES OF SILICON TETRACHLORIDE

Identification	Silicon Tetrachloride
Formula	sicl4
State (Std. Cond.)	Liquid
Molecular Weight, M	169.90
Boiling Point, Tb, °C	57.3
Melting Point, T_m , °C	-69.4
Critical Temp., T _C , °C	234.0
Critical Pressure, Pc, atm	37.0
Critical Volume, V _C , cm ³ /grmol	326.3
Critical Compressibility Factor, $\mathbf{Z}_{\mathcal{G}}$	0.290
Critical Density, ρ _C , gr/cm ³	0.5207
Acentric Factor, ω	0.2556

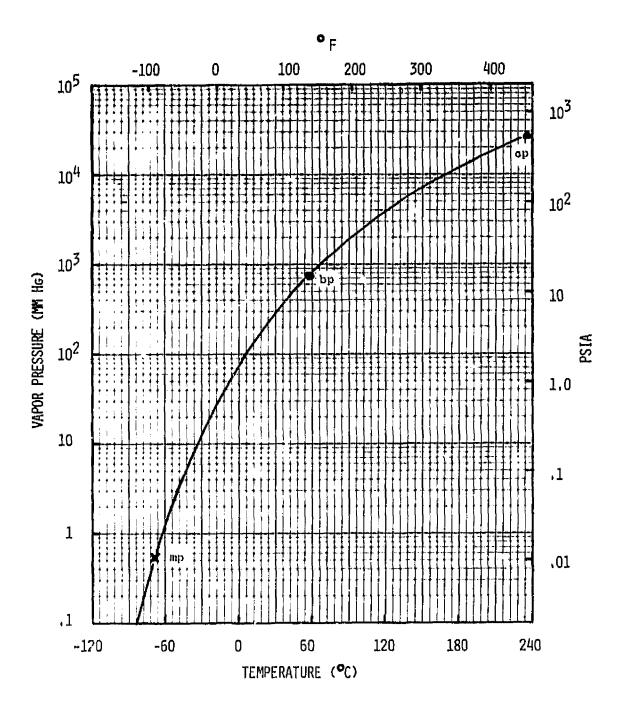


Figure 2.2-1 Vapor Pressure vs Temperature for Silicon Tetrachloride

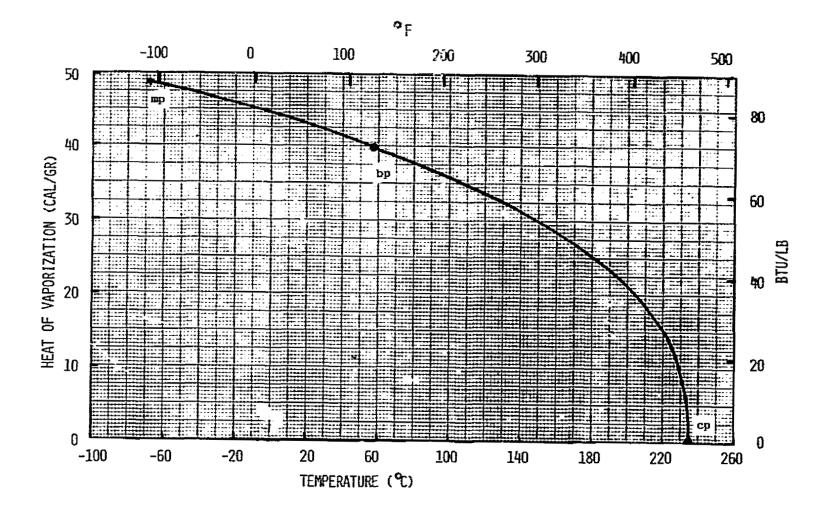


Figure 2.2-2 Heat of Vaporization vs Temperature for Silicon Tetrachloride

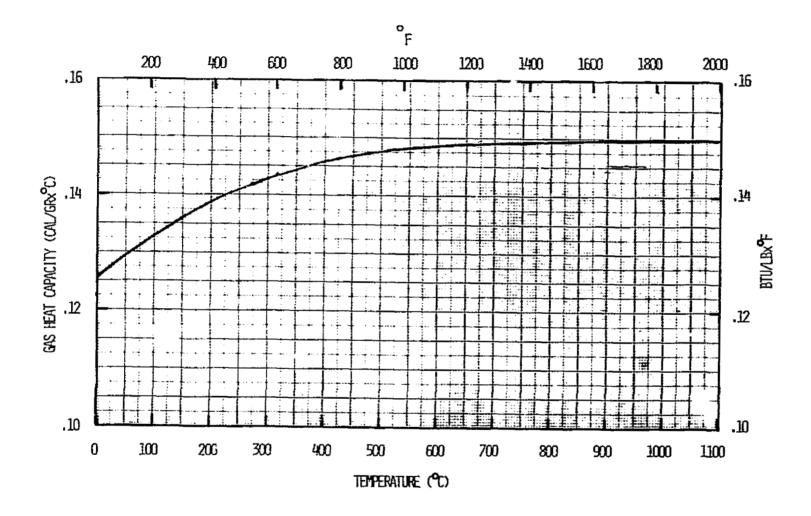


Figure 2.2-3 Gas Heat Capacity vs Temperature for Silicon Tetrachloride

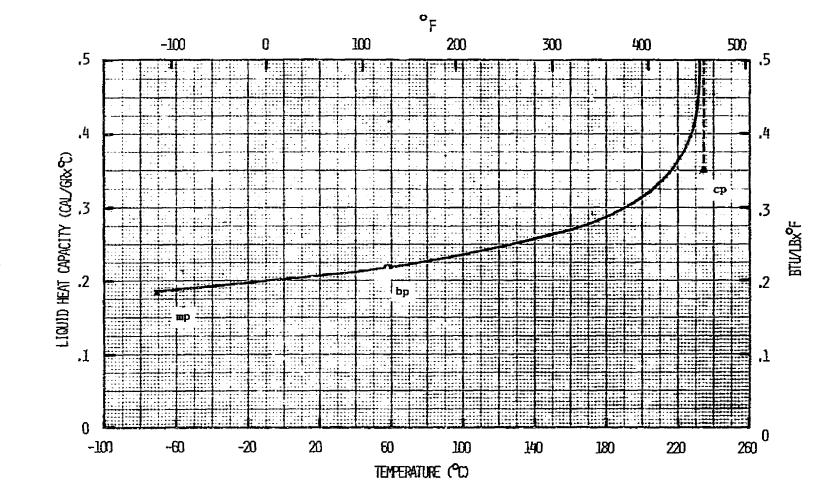


Figure 2.2-4 Liquid Heat Capacity vs Temperature for Silicon Tetrachloride

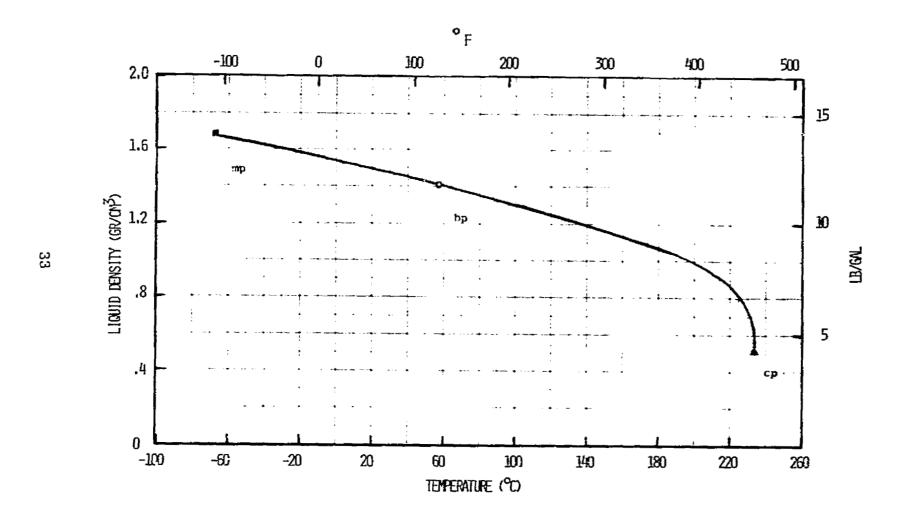


Figure 2.2-5 Liquid Density vs Temperature for Silicon Tetrachloride

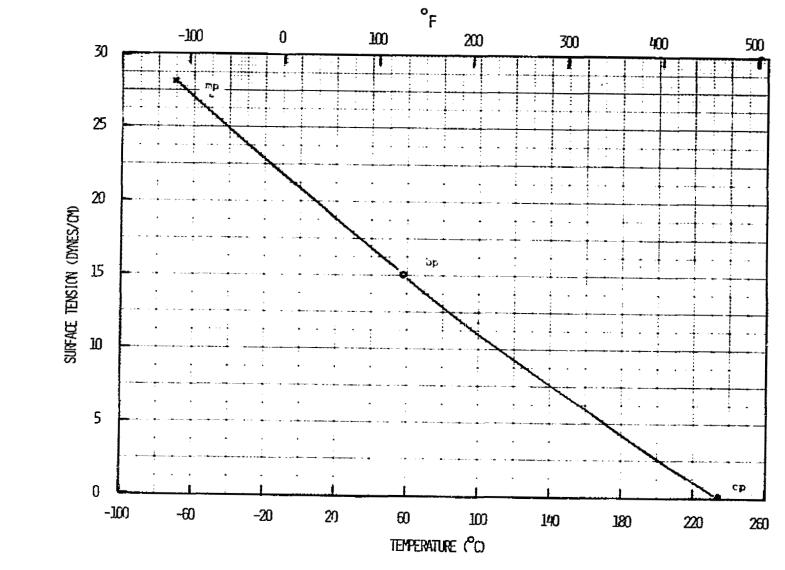


Figure 2.2-6 Surface Tension vs Temperature for Silicon Tetrachloride

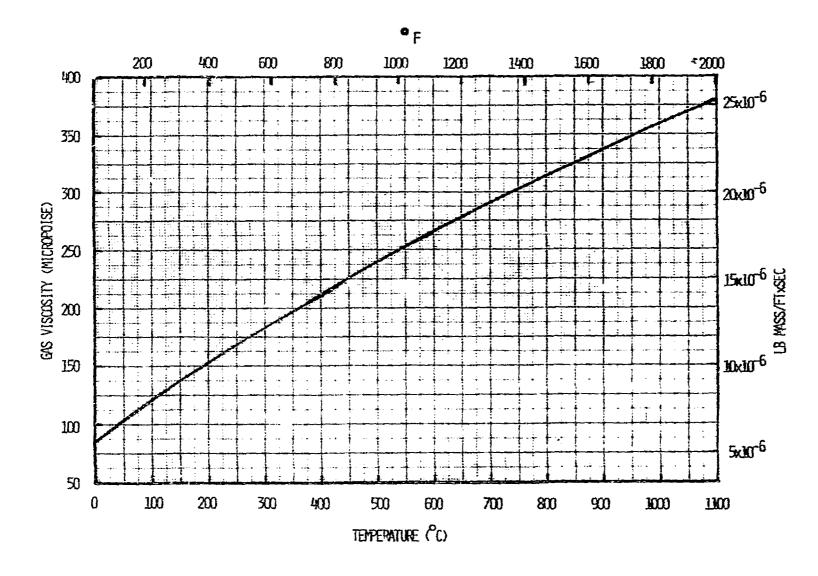


Figure 2.2-7 Gas Viscosity vs Temperature for Silicon Tetrachloride

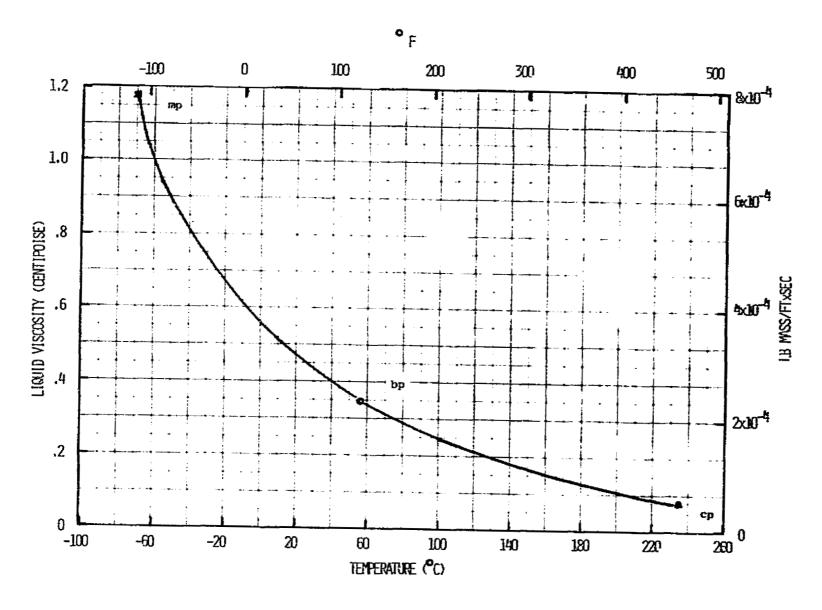


Figure 2.2-8 Liquid Visco y vs Temperature for Silicon Tetrachloride

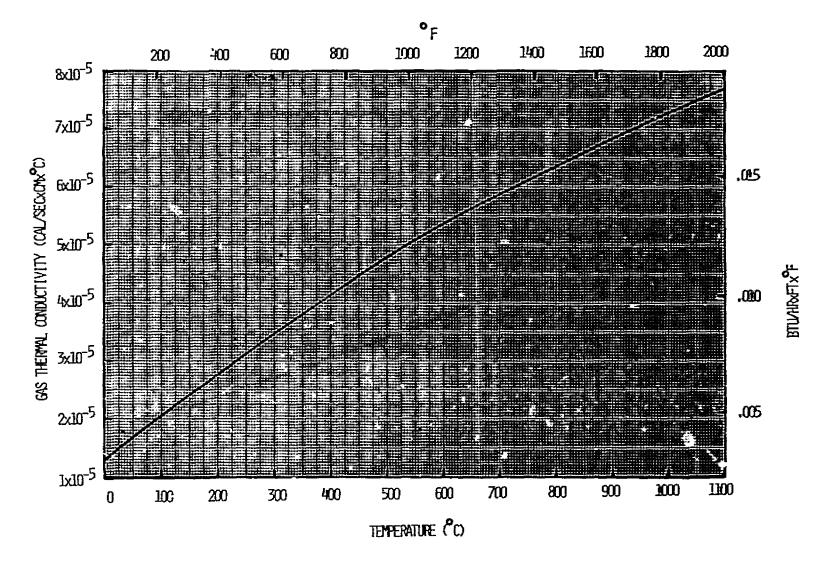


Figure 2.2-9 Gas Thermal Conductivity vs Temperature for Silicon Tetrachloride

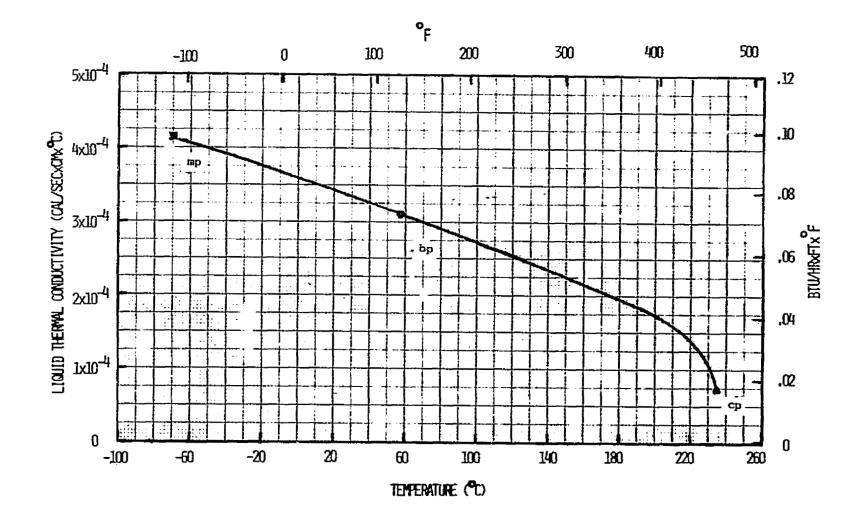


Figure 2.2-10 Liquid Thermal Conductivity vs Temperature for Silicon Tetrachloride

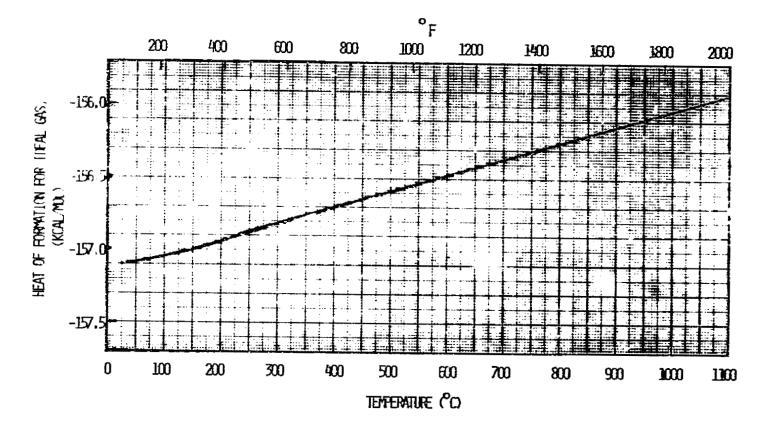


Figure 2.2-11 Heat of Formation vs Temperature for Silicon Tetrachloride

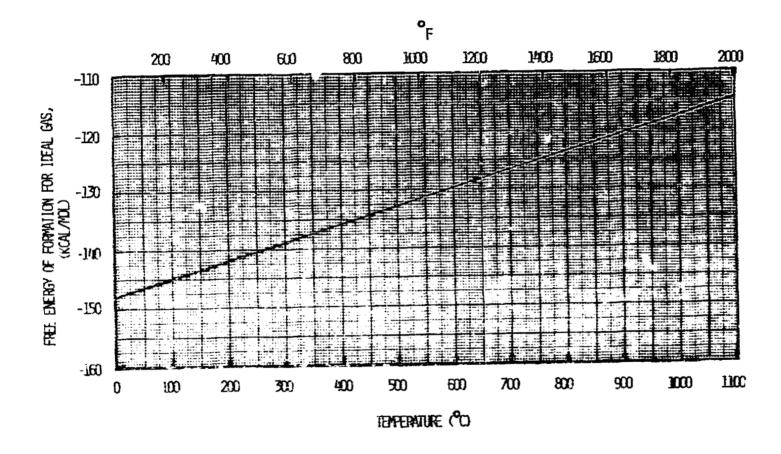


Figure 2.2-12 Free Energy of Formation vs Temperature for Silicon Tetrachloride

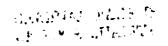
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2.3 Trichlorosilane Properties

Critical Properties (Table 2.3-1)

Experimental data for the critical temperature and critical volume are available (G33, G28) from a Russian investigation of orthobaric densities and critical paramaters. The critical pressure for trichlorosilane was estimated by Lydersen method (G62, G67):

$$P_{C} = \frac{M}{(\Sigma \Delta p + 0.34)^{2}}$$
 (2.3-1)

where $P_{\rm C}$ is critical pressure (atm), M is molecular weight (gr/gr-mol), and Δp is critical property increments for atoms making up the molecule. This method produced only 1.6% error when compared with the experimentally determined critical pressure of silicon tetrachloride.

The critical compressibility factor, Z_{c} , was calculated form its definition:

$$z_{c} = \frac{P_{c}V_{c}}{R T_{c}}$$
 (2.3-2)

Vapor Pressure (Figure 2.3-1)

Observed vapor pressure data from several sources (G15, G29, G44, G61) for trichlorosilane are in general agreement from -85°C to just above the boiling point. The experimental data were extended to cover the entire liquid phase using the YSSP correlation relation (G63):

$$\log P_V = A + \frac{B}{T} + C \log T + DT$$
 (2.3-3)

where

 P_v = vapor pressure of saturated liquid, mm of Hg

A, B, C, D = correlation constants for chemical compound

T = temperature, °K

The deviation of experimental and correlation results was small at 0.8% error for the 36 available data points.

Heat of Vaporization (Figure 2.3-2)

Heat of vaporization data for trichlorosilane are available only at the boiling point (G25, G18, G38, G46, G27). Using the known value at the boiling point, Watson's correlation (G52) was used to extend the heat of vaporization over the entire liquid phase:

$$\Delta H_{V} = \Delta H_{V1} \left[\begin{array}{c} T_{C} - T \\ T_{C} - T_{1} \end{array} \right]^{n}$$
 (2.3-4)

where n = .38 and $\Delta H_{\rm vl}$ applies at the boiling point (T₁).

Heat Capacity (Figures 2.3-3 and 2.3-4)

Heat capacity of the ideal gas at low pressure has been calculated by various Russian (G23, G25, G45, G11), American (G53, G56) and other (G6, G30) workers. The values, taken from various structural and spectral data, are in close agreement. The JANAF values (G53) were selected.

The liquid heat capacity of trichlorosilne is reported to be .23 between 25 and 60°C (G19, G46). The values are extended over all liquid temperatures by the relationship:

Heat Capacity x Density = Constant
$$(2.3-5)$$

The constant, C., was estimated to be 0.298.

Testing of this relaitonship with available data for silicon tetrachloride produced an average deviation of 4%.

Liquid Density (Figure 2.3-5)

Liquid density data for trichlorosilane are available from -10°C to the critical point (G33, G32, G61, G12, G26). The experimental data was extrapolated to the melting point by use of the Yaws-Shah relationship (G63) for saturated liquid:

$$\rho_{L} = AB^{-(1-T_{r})}^{2/7}$$
 (2.3-6)

where A=.4856 and B=.2618. Correlation values and experimental results were in close agreement. The deviation was less than 1% for the $\frac{31}{2}$ problems data points from several independent sources.

Surface Tension (Figure 2.3-6)

Data for the surface tension of trichlorosilane are available from 0°C to 40°C (G32, G26). These data were extended using the Othmer relations (G62):

$$\sigma = \sigma_1 \left[\frac{T_c - T}{T_c - T_1} \right]^n \tag{2.3-7}$$

where σ_1 = surface tension at T_1 , dynes/cm, and n = the correlation parameter, l.2. The other parameters have their usual meaning. Deviations between data and correlation values were 3% or less, largely due to the deviations between reported experimental values.

Viscosity (Figures 2.3-7 and 2.3-8)

Data for the gas viscosity of trichlorosilane were available only at 0° C and at boiling point (G25). The values at higher temperatures were estimated using the modified and revised corresponding-state method of Thodos and Yoon (G67, G68):

$$\eta_{c}\xi = 4.610 \text{ T}_{c}^{0.618} - 2.04 \text{ e}^{-0.-449}\text{T}_{r} + 1.94 \text{ e}^{-4.058}\text{T}_{r} + 0.1 \quad (2.3-8)$$

where $\eta_{\rm c}$ = viscosity, ξ = $T_{\rm c}^{-1/6} {\rm M}^{-1/2} {\rm P_c}^{-2/3}$, and $T_{\rm r}$ is the reduced temperature. The percentage error was less than .4%. Testing with silicon tetrachloride gave good agreement of correlation and experimental results (16 data points produced a 2% deviation).

Liquid viscosity data for trichlorosilane are available from -7°C to 60°C (G32, G26, G19, G25, G46). At low temperatures (from the boiling to the melting point), values were estimated using the log η_L vs 1/T linear relationship. At high temperatures (up to the critical point), the Stiel and Thodos correlation was used with $\mu_L \xi = f(Z_C, T_T)$ where $f(Z_C, T_T)$ is given as a generalized liquid v scosity correlation (G62). The percentage error with the available experimental data was about 2%.

Thermal Conductivity (Figures 2.3-9 and 2.3-10)

The gaseous thermal conductivity of trichlorosilane has recently been reported from 46°C to 350°C (G66). The experimental values were extended using a modified form of the Misic and Thodos correlation (G63, G67):

$$\lambda_{\rm G} = c_{\rm p}/\gamma \ (10^{-6}) (14.52 \ T_{\rm r} - 5.14)^{\rm n}$$
 (2.3-9)

where n = .71. The average absolute percentage error was 1.5%.

Liquid thermal conductivity data for trichlorosilane are not available. Using the estimation method of Sheffy and Johnson (G62):

$$\lambda_{T_{i}} = \{(4.66)(10^{-3})[1-.00126(T-T_{in})]/T_{in}^{2.16}M^{300}\}$$
 (2.3-10)

 $\lambda_{\rm L} = 2.783 \times 10^{-4}$ cal/cm x sec x °K was derived for the value at 60°C.

Using the Pachaiyappan-Vaidyanathan method of estimation (G64):

$$\lambda_{L} = 8.84 \times 10^{-4} c_{S} \rho_{L}$$
 (2.3-11)

the value of 2.64 \times 10⁻⁴ ca./cm \times sec \times 4K was derived for 60°C.

These estimation methods produced errors of 16% and 17.5%, respectively, on the one published value for ${\rm SiCl}_4$; and hence, should be taken to represent only an order of magnitude estimate. The estimate was extended over the entire liquid range using a modification of the Stiel and Thodos method (G62, G63):

$$\lambda_{L} = \frac{f(\rho_{r})}{\gamma Z_{c}^{5}} + \lambda_{G}$$
 (2.3-12)

Heat and Free Energy of Formation (Figures 2.3-11 and 2.3-12)

Values of the heat (ΔH_f) and Gibb's free energy of formation (ΔG_f) for the ideal gas are available from various Russian (G11, G45), American (G53) and other (G6, G30) sources and are in close agreement. The American values were selected.

TABLE 2.3-1
Critical Constants and Physical Properties of Trichlorosilane

Identification	Trichlorosilane
Formula	siHCl ₃
State (std. cond.)	liquid
Molecular Weight, M	135.453
Boi'ing Point, Tb, °C	31.8
Melting Point, T _m , *C	-126.6
Critical Temp., Tc, °C	206
Critical Pressure, P _C , atm	40.01*
Critical Volume, V _c , cm ³ /gr mol	268
Critical Compressibility Factor, Z	.273*
Critical Density, pc, gr/cm3	.505
Acentric Factor (Ω)	.188*

^{*}Estimated

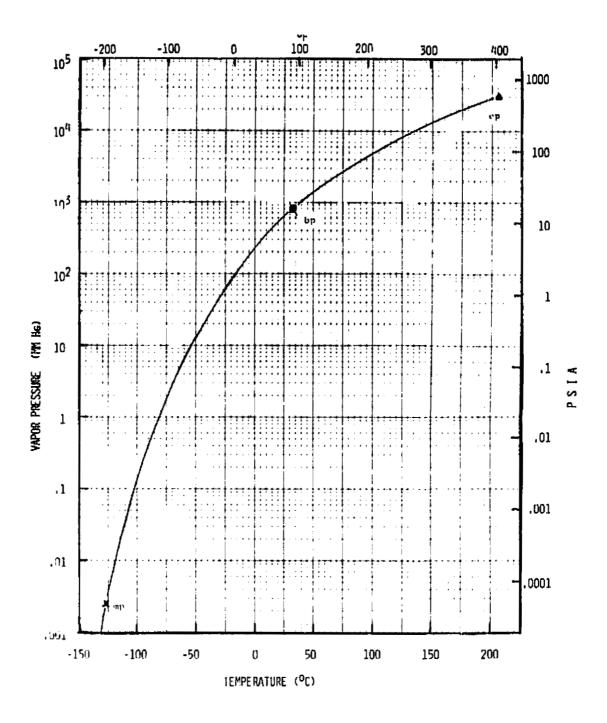


Figure 2.3-1 Vapor Pressure vs Temperature for Trichlorosilane



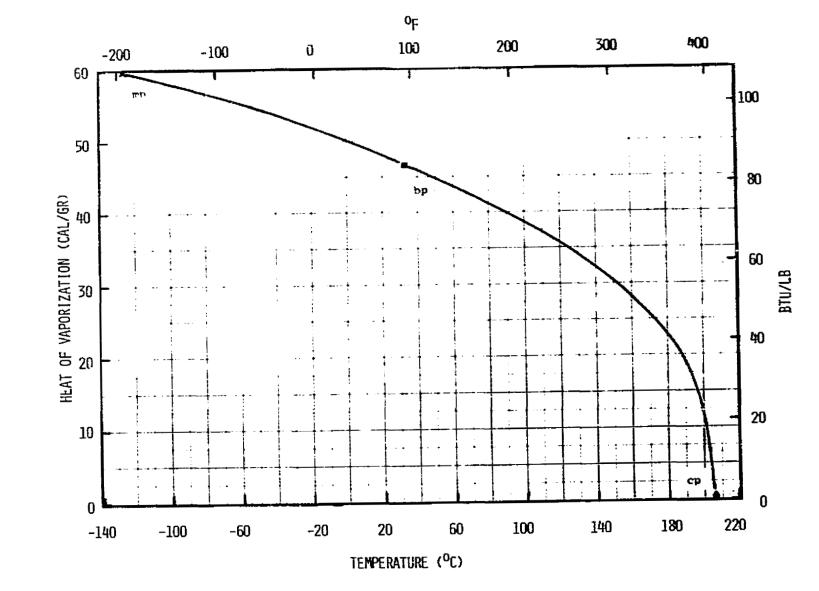


Figure 2.3-2 Heat of Vaporization vs Temperature for Trichlorosilane

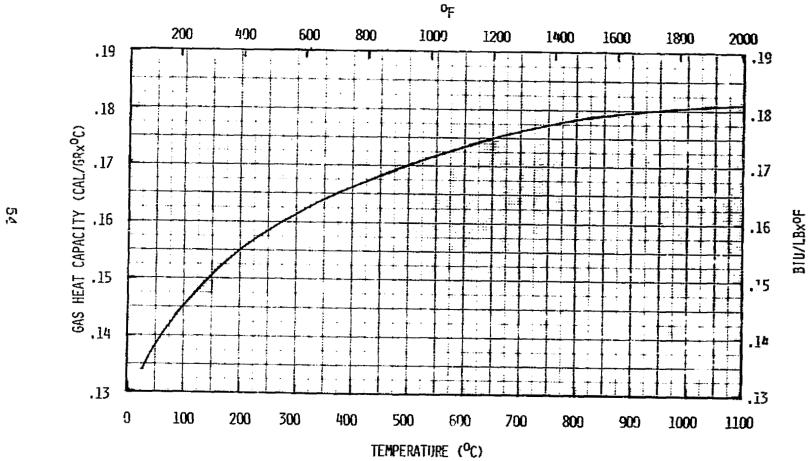


Figure 2.3-3 Gas Heat Capacity vs Temperature for Trichlorosilane

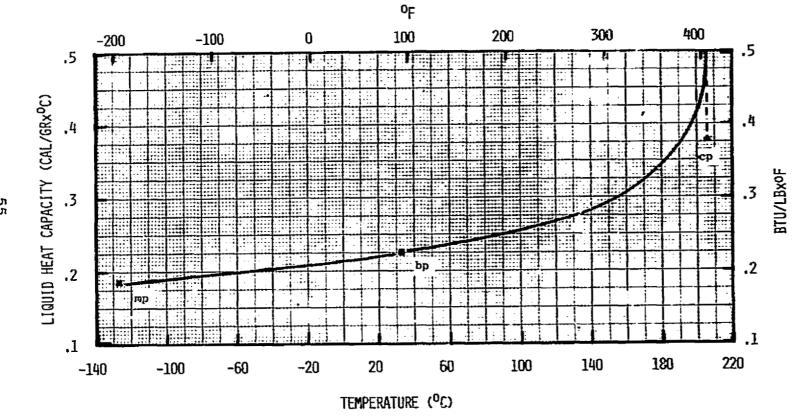


Figure 2.3-4 Liquid Heat Capacity vs Temperature for TrichlorosilaLe

Figure 2.3-5 Liquid Density vs Temperature for Trichlorosilane

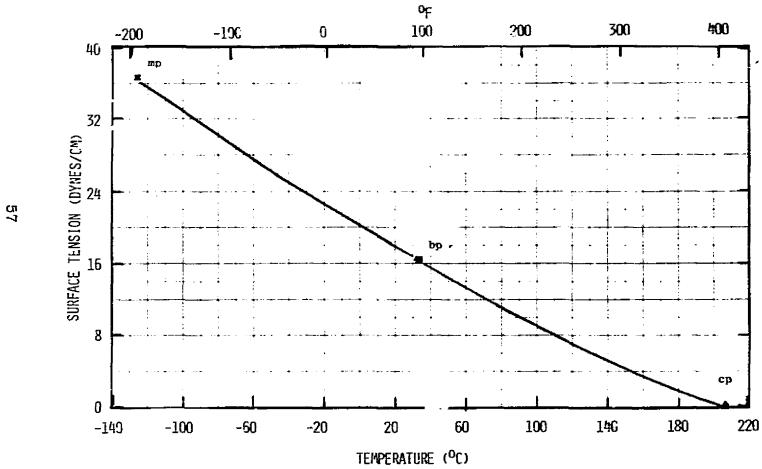


Figure 2.3-6 Surface Tension vs Temperature for Trichlorosilane

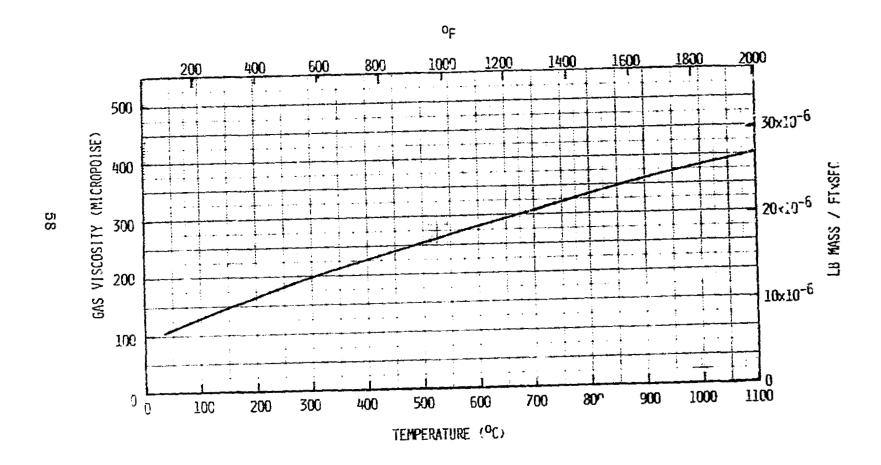


Figure 2.3-7 Gas Viscosity vs Temperature for Trichlorosilane

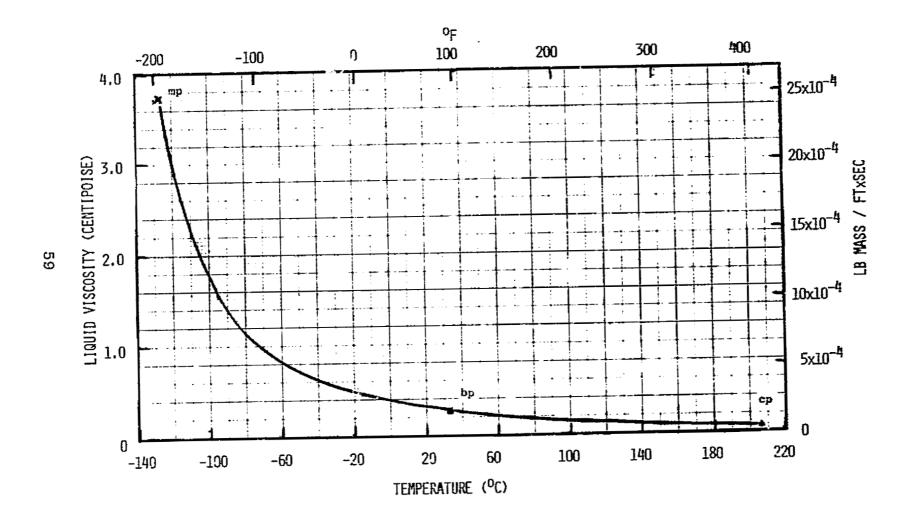


Figure 2.3-8 Liquid Viscosity vs Temperature for Trichlorosilane

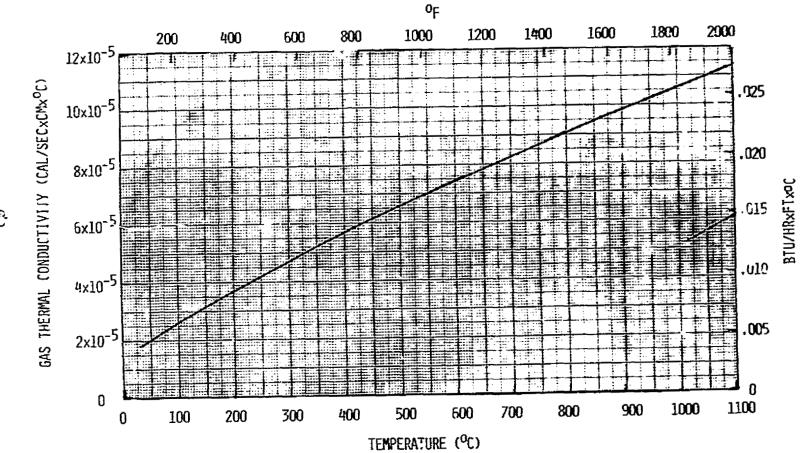


Figure 2.3-9 Gas Thermal Conductivity vs Temperature for Trichlorosilane

o_F

100

200

0

-200

-100

Figure 2.3-10 Liquid Thermal Conductivity vs Temperature for Trichlorosilane

TEMPERATURE (OC)

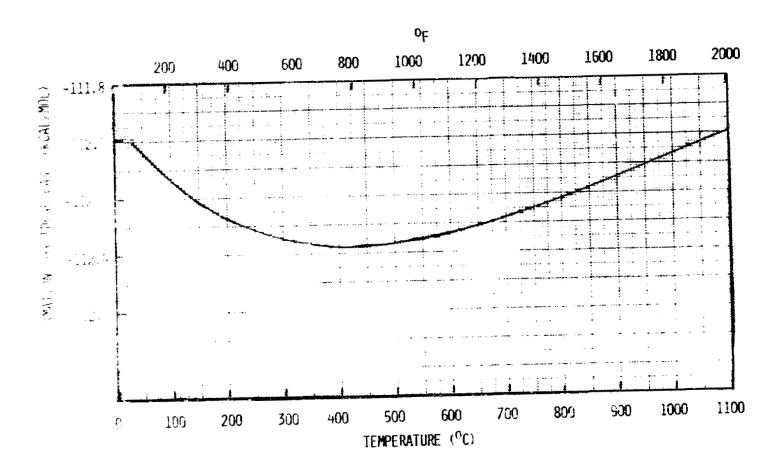


Figure 2.3-11 Heat of Formation vs Temperature for Trichlorosilane

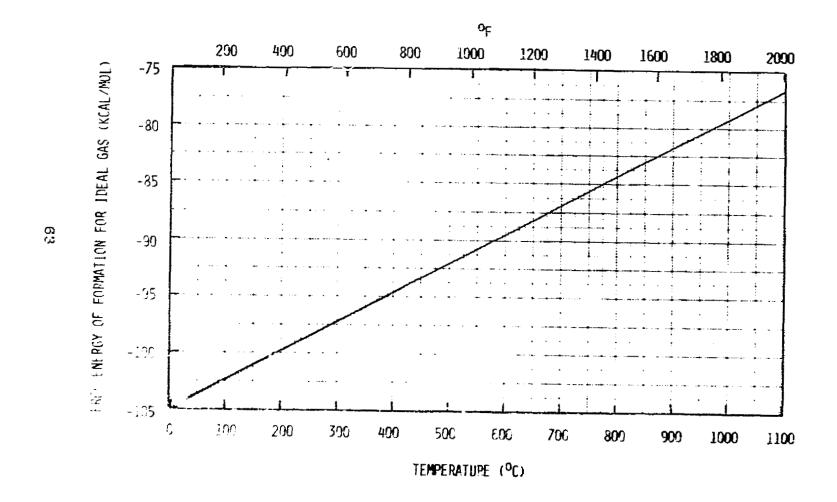


Figure 2.3-12 Free Energy of Formation vs Temperature for Trichlorosilane

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2.4 Dichlorosilane Properties

Physical Properties and Critical Constants (Table 2.4-1)

Physical properties and critical constants are listed in Table IA-1 for dichlorosilane. Values of critical temperature, $T_{\rm C}$, critical pressure, $P_{\rm C}$, and critical volume, $V_{\rm C}$, for dichlorosilane were estimated by using Lydersen's structural contribution method with derived critical preparty increments for silicon (H16). This method produced only 2.3% error for $T_{\rm C}$ and 3.4% error for $V_{\rm C}$ when compared with the experimental values of trichlorosilane and it produced 0% error for $T_{\rm C}$, $V_{\rm C}$, and $P_{\rm C}$ when compared with the known values of silicon tetrachloride. The estimated values for the known values for the critical properties are also within reasonable agraement (4% for $T_{\rm C}$, 0.2% for $P_{\rm C}$, and 14% for $V_{\rm C}$) of calculated Russian values (H10).

The critical compressibility factor, $\mathbf{Z}_{\mathbf{C}}$, was determined from its definition:

$$z_{c} = \frac{V_{c}RT_{c}}{P_{c}}$$
 (2.4-1)

The result from Eq. (IA-1) was the same as that derived by the Garcia-Barcena' boiling point method (H16):

$$Z_{c} = f(T_{b}) - g(T_{b}/M)$$
 (2.4-2)

Vapor Pressure (Figure 2.4-1)

The vapor pressure of dichlorosilane has been determined from -80°C to 30°C (H23, H35). The experimental data was extended over the entire liquid range using the YSSP vapor pressure correlation (H30):

$$\log P_V = A + \frac{B}{m} + C \log T + DT + ET^2$$
 (2.4-3)

where P_V is the vapor pressure of saturated liquid, mm Hg; T is temperature, $^{\circ}K$; and A, B, C, D, E are correlation constants derived using a generalized least squares computer program. Average absolute deviation was about 1% for the 13 experimental data points.

Heat of Vaporization (Figure 2.4-2)

Heat of vaporization data for dichlorosilane are available only at the boiling point (H1, H0, H10, H19, H31). Using the known value at the boiling point, Watson's correlation was used to extend the heat of vaporization over the entire liquid phase:

$$\Delta H_{\mathbf{V}} = \Delta H_{\mathbf{V}_1} \left[\frac{T_{\mathbf{C}} + T}{T_{\mathbf{C}} - T_1} \right]^{n} \tag{2.4-4}$$

where n = .38 and ΔH_{V_1} applies at the boiling point (T_1) .

Heat Capacity (Figures 2.4-3 and 2.4-4)

Ideal gas heat capacity data for dichlorosilane are available from various American (H5, H13, H25, H26), Russian (H6, H7, H10, H12, H32) and other (N9, H33) workers. The vlaues, which are in close agreement, are based on bond additivities and spectral measurement. The JANAF values were selected.

Measured saturated-liquid heat capacity data for ...ichlorosilane are unavailable in the literature. Values were estimated from ~60°C to 60°C using the Yuan and Stiel corresponding state method (H16). For polar liquids, the correlation takes the form:

$$c_{\sigma_1} - c_p^0 = \Lambda c^{(0p)} + \omega (\Lambda c_{\sigma}^{(1p)} + x (\Lambda c_{\sigma}^{(2p)} + x^2 (\Lambda c_{\sigma}^{(3p)} + \omega^2 (\Lambda c_{\sigma}^{(4p)} + x\omega) (\Lambda c_{\sigma}^{(5p)})$$
(2.4-5)

where CB is the ideal gas heat capacity, ω is the acentric factor, X is the Stiel polar factor and the functions: $(AC_{C}^{-}(Op))$, etc. are tabulated as functions of the reduced temperature. The relationship that heat capacity times density is constant was used to extend the values over the entire liquid range. Application of the Yuan and Stiel correlations to silicon tetrachloride, trichlorosilane, and silicon tetrafluoride gave average absolute percentage errors of 3.1, 6.7, and 4.3 respectively. Due to the limited experimental data points, the calculated liquid heat capacities should be considered as order-of-magnitude estimates.

Density (Figure 2.4-5)

Liquid density data are available at the molti- τ point (H8, H9, H10, H18, H27) and at 7°C (H35). The limited data we extended over the entire liquid range using a modification of the Rackett equation:

$$\rho = \rho_{\rm c} z^{-(1-T_{\rm r})}^{2/7} \tag{2.4-6}$$

where $\rho_{\rm C}$ is critical density, Tr is reduced temperature and Z is a parameter defined by the experimental data.

Surface Tension (Figure 2.4-6)

The Brock and Bird corresponding states method (H16) was used to estimate the surface tension of dichlorosilane since no experimental data is available. The equation is:

$$\sigma = P_c^{2/3} T_c^{1/3} (0.133 \alpha_c - 0.281) (1 - T_r)^{11/9}$$
 (2.4-7)

where σ is surface tension, dynes/cm, $\alpha_{\rm C}$ is the Riedel parameter, $P_{\rm C}$ is critical pressure, atm.; $T_{\rm C}$ is critical tengerature, ${}^{\rm o}{\rm K}$; and $T_{\rm T}$ is the reduced temperature. Application of this method to bilton tetrachloride and trichlorosilane gave results within 4% and 0.8% absolute deviation with expensional data, respectively.

Viscosity (Figures 2.4-7 and 2.4-8)

Gas viscosity calculations at low pressure were made using the methods of (!) Youn and Thodos for non-hydrogen-bouding polar gases, (2) Golubev, and (3) Reichenberg (H16). Since the calculated values were in close agreement, they were fitted to the series expansion:

$$\eta_G = A + BT + CT^2$$
 (2.4-8)

where η_G is in micropoise; T is temperature, ${}^{\circ}K$; and A, B and C are computer derived parameters using a generalized least squares program. The average absolute percentage deviation was less than 1.8%.

Liquid viscosities at temperatures below the boiling point were calculated using the methods of Thomas, and of Morris (up to 60°C) (H16). Values from the boiling point to the critical point were calculated using the correlation methods of Leton and Stiel, and Stiel and Thodos (H16). Calculated values were extended over entire liquid range and fitted to the equation:

$$\log \eta_{T} = A + \frac{B}{T} + CT + DT^2$$
 (2.4-9)

where η_L is in centipoise; T is temperature, °K; and A, B, D and D are derived parameters using a generalized least squares computer program. This was done in order to fit together the calculated values which apply in the different temperature ranges. The average percentage deviation was 3.3% with the greater deviation being near the melting point; therefore, this should be considered to be an order-of-magnitude correlation.

Thermal Conductivity (Figures 2.4-9 and 2.4-10)

Gas-phase thermal conductivity data are available from 28°C to 350°C (H28). The data were correlated and extended to higher temperatures by a series expansion in temperature:

$$\lambda_{\rm G} = A + BT + CT^2 + DT^3$$
 (2.4-10)

where λ_G is gas thermal conductivity, cal/cm x sec x °C; T is temperature, °K; and A, B, C and D are computer derived constants characteristic of the chemical compound. The absolute deviation between data and correlation values was less than 0.5%.

Thermal conductivity data of the liquid phase is unavailable. Modifications of the estimation methods of Sato and Reidel (H16):

$$\lambda_{\rm L} = \frac{2.64 \times 10^{-3}}{{\rm M}^{1/2}} \frac{3 + 20 (1 - {\rm T_r})^{2/3}}{3 + 20 (1 - {\rm T_r})^{2/3}}$$
 (2.4-11)

and of Robbins and Kingrea (H15):

$$\lambda_{\rm L} = \frac{(88 - 4.94 \text{ H}) \times 10^{-3}}{\Delta S^*} \frac{.55}{T_{\rm w}} C_{\rm p_{\rm L}} \rho^{4/3}$$
 (2.4-12)

where used to derive values at 32°C. These modified estimation methods produced error of less than 1% absolute deviation on the one published value of SiCl₄. The average of the estimate at 32°C was extended over the entire liquid range using a modification of the Stiel and Thodos method (H16):

$$\lambda_{\rm L} = \frac{f(\rho_{\rm r})}{\Gamma z_{\rm c}^{5}} + \lambda_{\rm G}$$
 (2.4-13)

The modified Sato-Reidel equation produced a similar range of values. Since assumptions in these calculations include the accuracy of the one data point for silicon tetrachloride and the chemical similarities in a homologous series, these values should be considered only order of-magnitude estimates.

Heat and Free Energy of Formation (Figures 2.4-11 and 2.4-12)

Heat of formation and Gibb's free energy of formation for the ideal gas have been estimated by Russian (H32, H36) and American (H25) workers up to at least 1500°K. Some estimated values differ significantly having about 35% deviation for ΔH_f and about 45% deviation for ΔG_f (H32, H36). The JANAF values (H25) were selected.

TABLE 2.4-1

CRITICAL CONSTANTS AND PHYSICAL PROPERTIES OF DICHLOROSILANE

Identification	<u> Michlorosilane</u>
Formula	SiH2Cl2
State (std. cond.)	gas
Molecular weight, M	101.008
Boiling Point, Tb, °C	8.3
Melting Point, Tm, °C	-122.0
Critical Temperature, TC, °C	178.9*
Critical Pressure, Pc, atm	44.0*
Critical Volume, V _C , cm ³ /gr mol	228.3*
Critical Compressibility Factor, Zc	.276*
Critical Density, pc, gr/cm3	.4424*
Acentric Factor (w)	.1107

^{*}Estimated

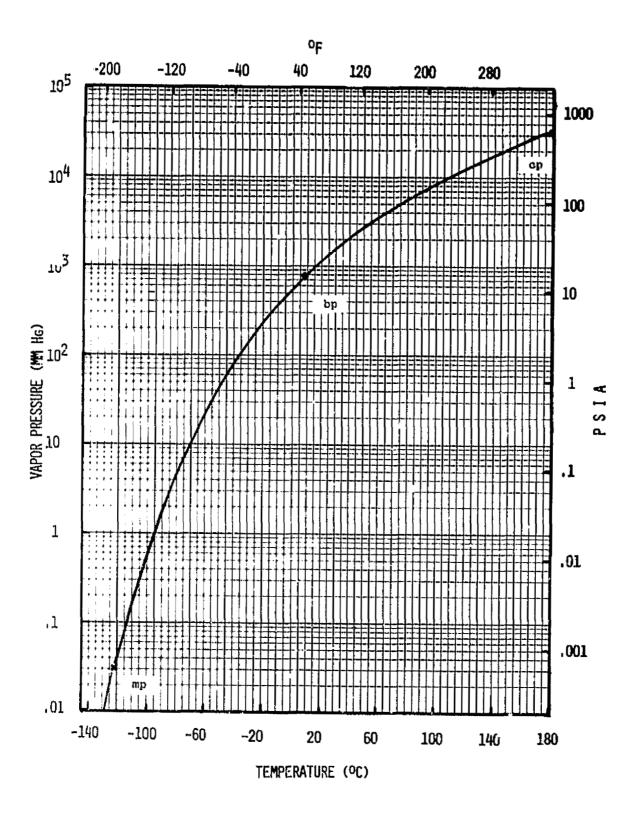


Figure 2.4-1 Vapor Pressure vs Temperature for Dichlorosilane

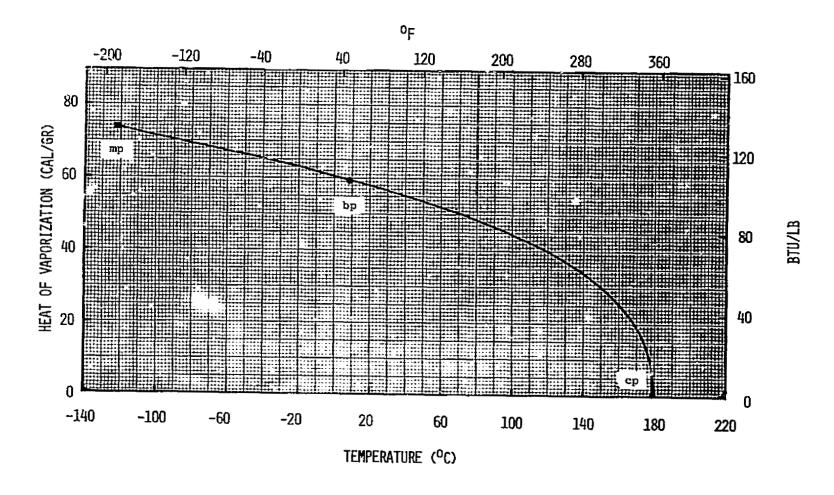


Figure 2.4-2 Heat of Vaporization vs Temperature for Dichlorosilane

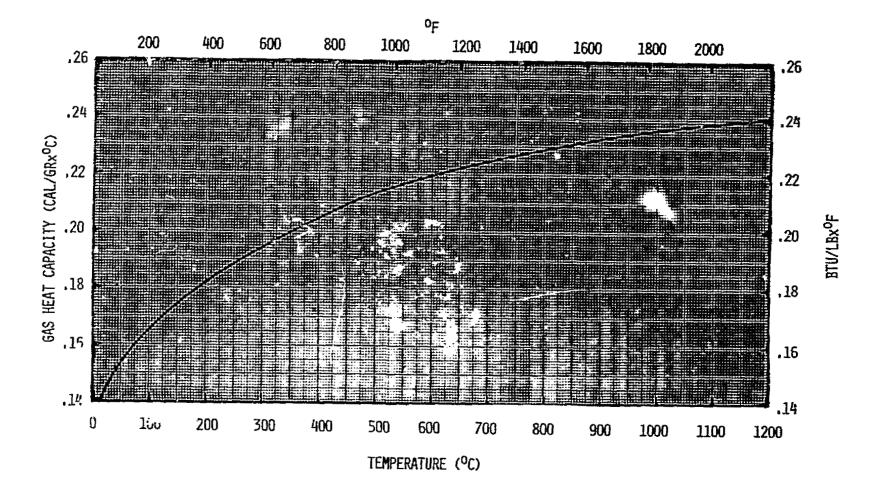


Figure 2.4-3 Gas Heat Capacity vs Temperature for Dichlorosilane

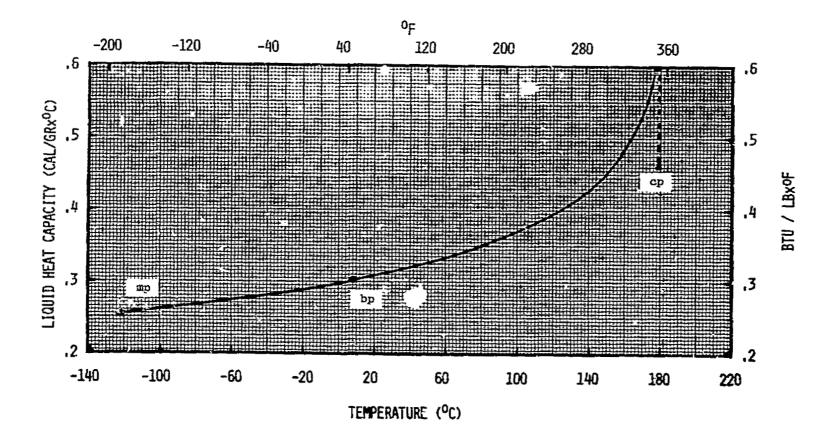


Figure 2.4-4 Liquid Heat Capacity vs Temperature for Dicklorosilane

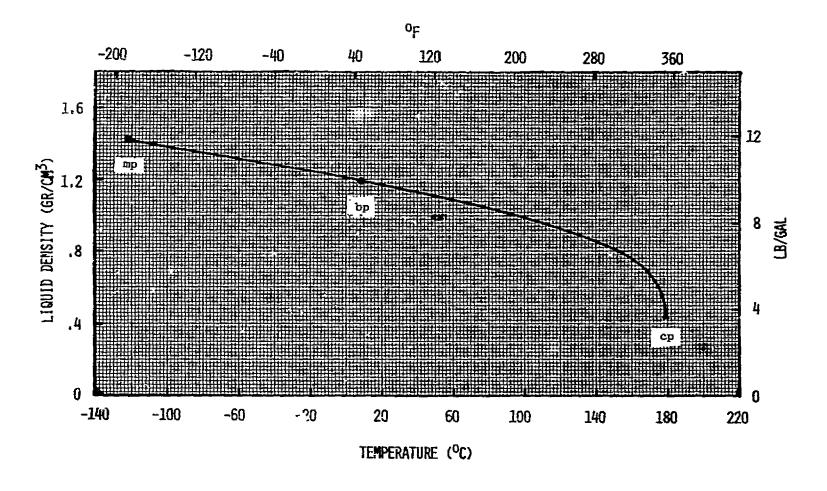


Figure 2.4-5 Liquid Density vs Temperature for Dichlorosilane

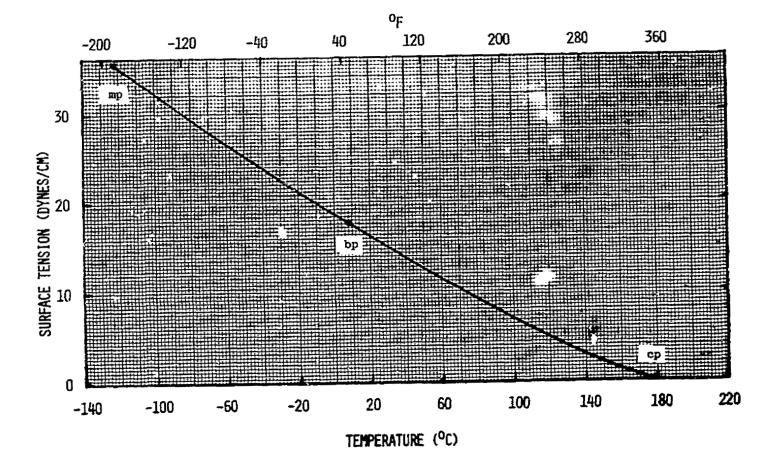


Figure 2.4-6 Surface Tension vs Temperature for Dichlorosilane

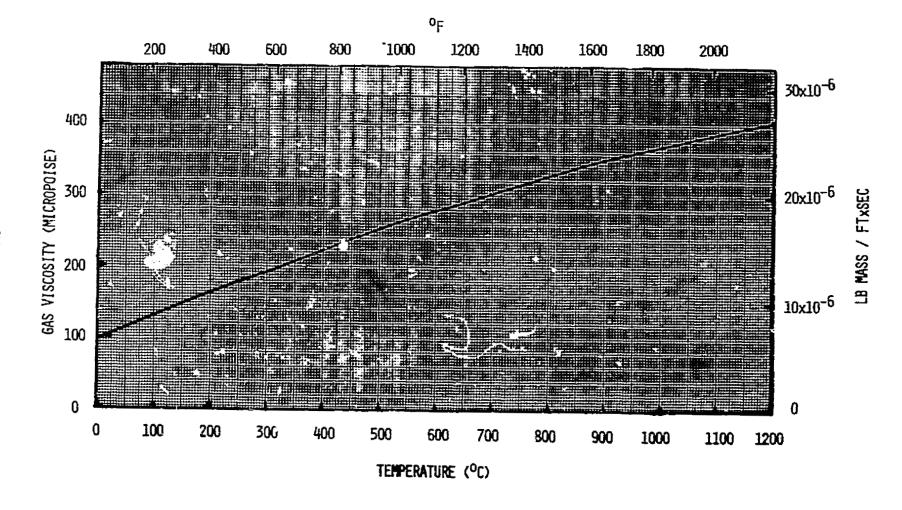


Figure 2.4-7 Gas Viscosity vs Temperature for Dichlorosilane

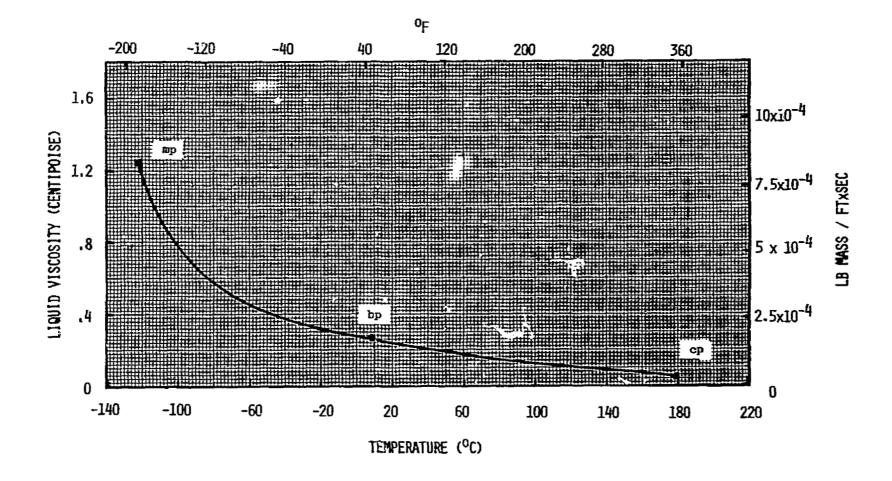


Figure 2.4-8 Liquid Viscosity vs Temperature for Dichlorosilane

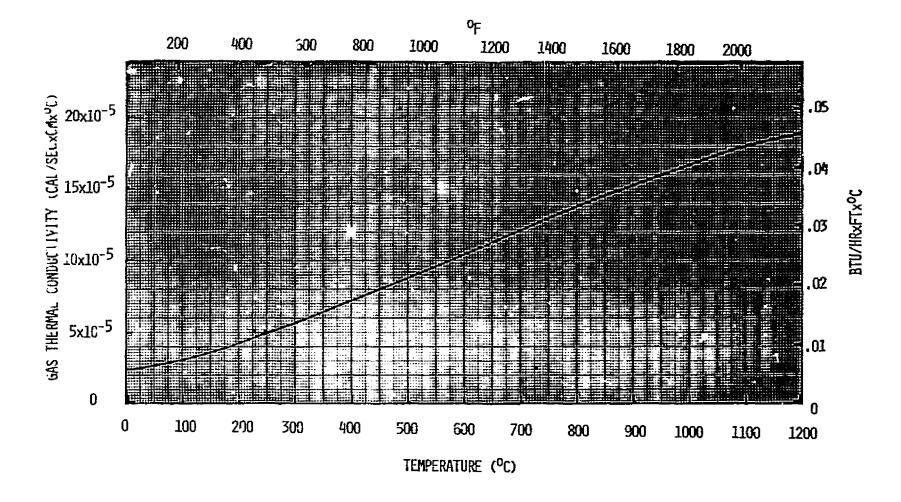


Figure 2.4-9 Gas Thermal Conductivity vs Temperature for Dichlorosilane

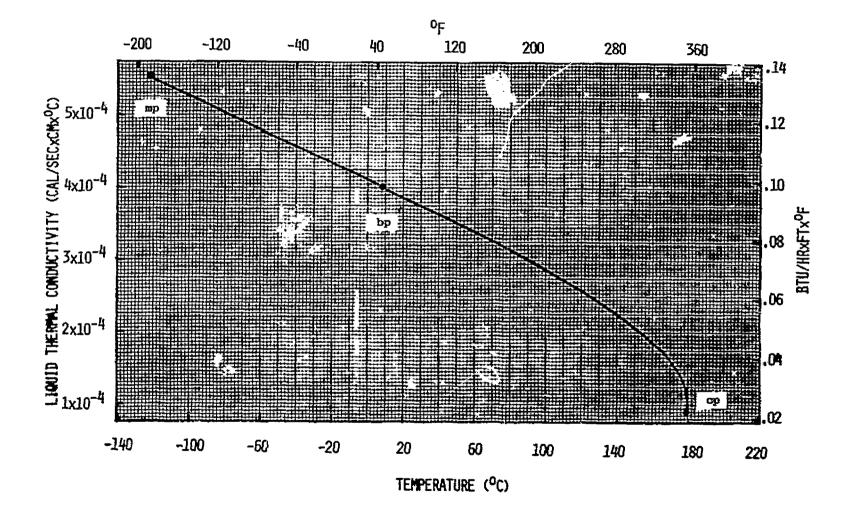


Figure 2.4-10 Liquid Thermal Conductivity vs Temperature for Dichlorosilane

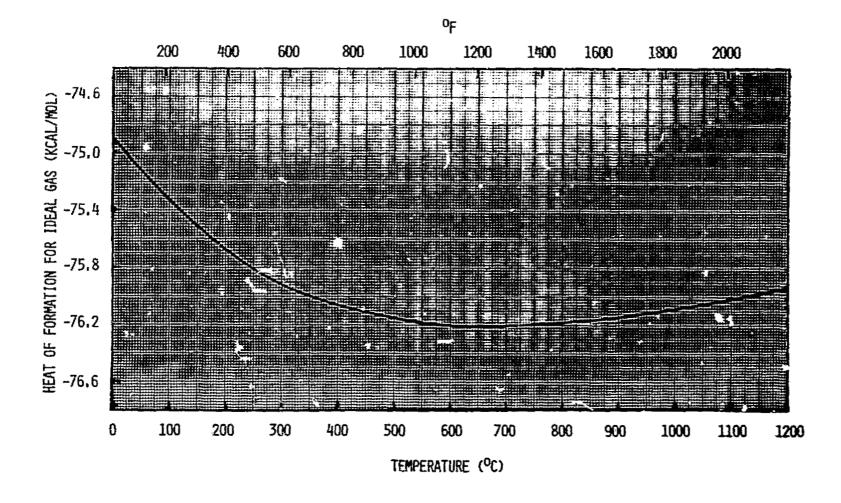


Figure 2.4-11 Heat of Formation vs Temperature for Dichlorosilane

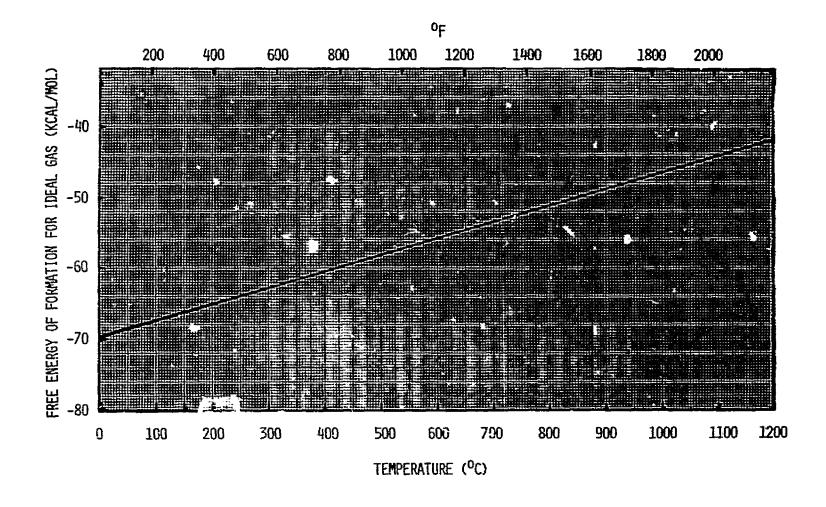


Figure 2.4-12 Free Energy of Formation vs Temperature for Dichlorosilane

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2.5 Silicon Tetrafluoride Properties

Physical Properties and Critical Constants (Table 3.5-1)

Physical properties are listed in Table IA-1 including the boiling point (sublimation temperature, where the vapor pressure of the solid is 760 mm Hg.). Two temperature are given in the literature for the melting point (triple point) of silicon tetrafluoride based on the work of Patnode and Papish (F37) in 1930 and the work of Pace and Mosser (F36) in 1963. The more recent work was selected as the melting point (triple point, where solid, liquid and vapor are in equilibrium).

Experimental data for the critical temperature and critical pressure of silicon tetrafluoride have been determined (F4). The critical compressibility factor, $\mathbf{Z}_{\mathbf{C}}$, was estimated from the Garcia'-Barcena' correlation (F39):

$$Z_{c} = f(T_{b}) - g(T_{b}/M)$$
 (2.5-1)

where $T_{\rm b}$ is the normal boiling point, $^{\rm e}$ K, and M is the molecular weight. When this method was applied to sulfur hexafluoride, another subliming inorganic fluoride, the calculated value of the critical compressibility factor only deviated 3.3% from the known value. From the estimated value of $Z_{\rm c}$, the critical volume, $V_{\rm c}$, was calculated by the rearrangement of the definition of $Z_{\rm c}$.

$$V_{c} = \frac{Z_{c}RT_{c}}{P_{c}}$$
 (2.5-2)

Vapor Pressure (Figure 2.5-1)

The vapor pressure of silicon tetrafluoride has been determined experimentally from 50° below the sublimation point to near the critical point (F4, F36, F37, F41). The experimental data were extended to cover the liquid range (from triple point to critical point), and for the solid (below the triple point using the YSSP correlation in each case:

$$\log P_{V} = A + \frac{B}{T} + C \log T + DT \qquad (2.5-3)$$

At the higher temperatures the deviation of experimental and correlation results are 2% while the average percent error was 4.6% in the lower temperature range. Much of the deviation is due to rather poor agreement in the literature for the experimenal values.

Heat of Vaporization (Figure 2.5-2)

Heat of vaporisation data for silicon tetrafluoride are reported near the triple point (F24, F29, F36, F41, F60). Using the selected value (F36), Watson's correlation (F39) was used to extend the heat of vaporisation over the entire liquid phase:

$$\Delta H_{V} = \Delta H_{V_{1}} \left[\frac{T_{c} - T}{T_{c} - T_{1}} \right]^{n}$$
(2.5-4)

where n=.38 and T_1 is the boiling point. The value of H_{V_1} was effectively confirmed using the Clausius-Clapyron equation and literature vapor pressure values (F36).

heat capacity (Figures 2.5-3 and 2.5-4)

Heat capacity of the ideal gas at low pressure has been calculated from 0°C to 1200°C (F48, F63). These values, including other values covering smaller temperature ranges (F11, F24, F26, F29, F35, F60), were taken from various structural and spectral data and are in close argeoment. The JANAF values (F48) were selected.

The liquid heat capacity of silicon tetrafluoride is reported near the sublimation point (F36). The values are extended over all liquid temperatures by the relationship:

The estimated constant was 0.473. Testing of this relationship with available data (4 data points) for silicon tetrafluoride produced an average absolute deviation of 0.8% error.

Liquid Donsity (Figure 2.5-6)

taquid density data for silicon tetrafluoride are available only within about 30 degrees of the triple point (F26, F60). The experimental data were extrapolated to the critical point by use of a modification of the Rackett equation (F65, F66):

$$\rho = \rho_c z^{-(1-T_r)}^{2/7}$$
 (2.5-6)

where $r_{\rm c}$ is the critical density, $T_{\rm r}$ is the reduced temperature, and Z is a parameter derived from available data. Comparison of the calculated and experimental values of 5 data points gave 0.57% average absolute error.

Surface Tension (Figure 2.5-6)

Experimental data are not available for the surface tension of silicon tetrafluoride. The Brock and Bird corresponding states technique was used to estimate the surface tension (F39):

$$\alpha = P_c^{2/3} T_c^{1/3} (0.153 \alpha_c - 0.281) (1-T_r)^{11/9}$$
 (2.5-7)

where α is surface tension, dynes/cm; α_{c} is the Riedel parameter; P_{c} is critical pressure, atm; T_{c} is the critical temperature, $^{\circ}K$; and T_{r} is the reduced temperature. Application of this technique to silicon tetrachloride and sulfur hexafluoride gave results within 4% and 1% absolute deviation with experimental data, respectively.

Viscosity (Figures 2.5-7 and 2.5-8)

Experimental data for the gas viscosity of silicon tetrafluoride are available from about room temperature to above 300°C (F13, F32). The values at higher temperatures were estimated using the relationship:

$$\log \eta_{\rm G} = A + BT + CT^2$$
 (2.5-8)

The average absolute percentage error was 1.74% when correlated values were compared with the 28 experimental data points.

No experimental data are available for the liquid viscosity of silicon tetrafluoride. Estimates were derived applying the Letsou-Stiel high-temperature liquid-viscosity correlation (F65):

$$\eta_{L} \xi = (\eta_{L} \xi)^{O} + \omega (\eta_{L} \xi)^{1}$$
 (2.5-9)

where the parameters $(\eta_L \xi)^0$ and $(\eta_L \xi)^1$ are functions of reduced temperature, ω is the acentric factor and $\xi = T_c^{1/6}/M^{1/2}p_c^{2/3}$. This correlations gave results within 17% and 48% absolute deviaiton for the experimental values of silicon tetracloride and sulfur hexafluoride, respectively. Since liquid viscosity estimation methods may be grossly inaccurate (F69), these values must be assumed to be order of magnitude estimates only.

Thermal Conductivity (Figures 2.5-9 and 2.5-10)

The gaseous thermal conductivity of silicon tetrafluoride has been reported from about room temperature to 350°C (F9, F68). The experimental values were extended using a modified form of the Misic and Thodos correlation (F66):

$$\lambda_{\rm G} = \frac{C_{\rm p}}{7} (10^{-6}) (14.52T_{\rm r} - 5.14)^{\rm n}$$
 (2.5-10)

where n = .63, Γ is $T_C^{-1/6} M^{1/2}/P_C^{-2/3}$, T_r is the reduced temperature and C_p is the gaseous heat capacity. The average percentage error was less than one percent.

No liquid thermal conductivity data are available; however, values were estimated using the Sato-Riedel equation (65):

$$\lambda_{L} = \frac{2.64 \times 10^{-3}}{M^{1/2}} \frac{3 + 20(1-T_{r})^{2/3}}{3 + 20(1-T_{r_{b}})^{2/3}}$$
(2.5-11)

where M is molecular weight, T_r is the reduced temperature, and $T_{r_{||}}$ is the reduced temperature at the boiling point. This correlation gave 34% error with the single experimental data point for silicon tetrachloride and 24% error for sulfur hexafluoride with the several experimental data points. There is considerable deviation of values among the several different data sources (22% maximum deviation). The present results should be taken only to represent an order of magnitude estimate.

Heat and Free Energy of Formation (Figures 2.5-11 and 2.5-12)

Many American workers (F20, F24, F29, F34, F35, F48, F57, F58) and others (F40, F43) have reported heats of formation was well as Gibb's free energy of formation (F48, F56, F58) for the ideal gas. The JANAF values (F48) were selected.

TABLE 2.5-1

PHYSICAL PROPERTIES AND CRITICAL CONSTANTS OF SILICON TETRAFLUORIDE

<u>Identification</u>	Silicon Tetrafluoride
Formula	SiF ₄
State (std. cond.)	gas (colorless)
Molecular weight, M	104.08
Boiling point, T _b , °C (sublimation point)	-95.7 (760 mm Hg)
Melting Point, T _m , °C (triple point)	-86.8 (1679 mm Hg) *Ref. F36 -90.2 (1318 mm Hg) Ref. F37
Critical Temp, Tc, °C	-14.15
Critical Pressure, Pc, atm	36.66
Critical Volume, Vc, cm3/gr mol	165**
Critical Compressibility Factor, Z	0.284**
Critical Density, p _c . gr/cm ³	0.6308**
Acentric Factor (0)	0.4086

^{*} Selected Value

^{**}Estimated

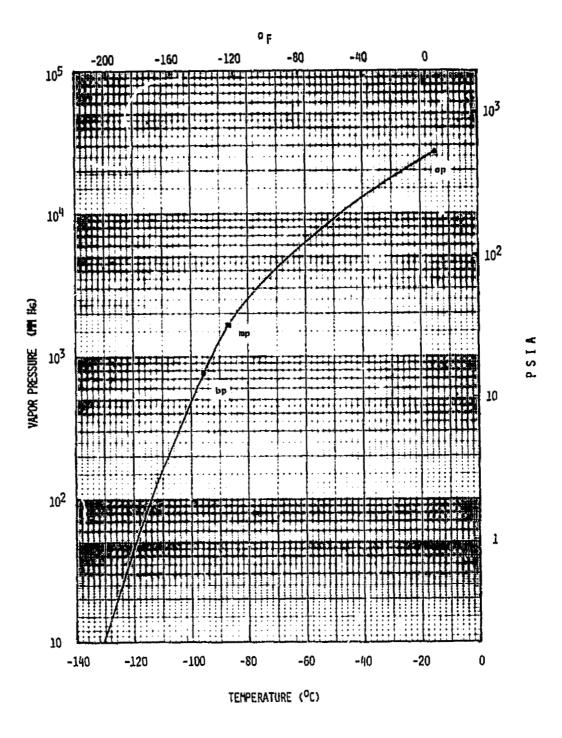


Figure 2.5-1 Vapor Pressure vs Temperature for Silicon Tetrafluoride

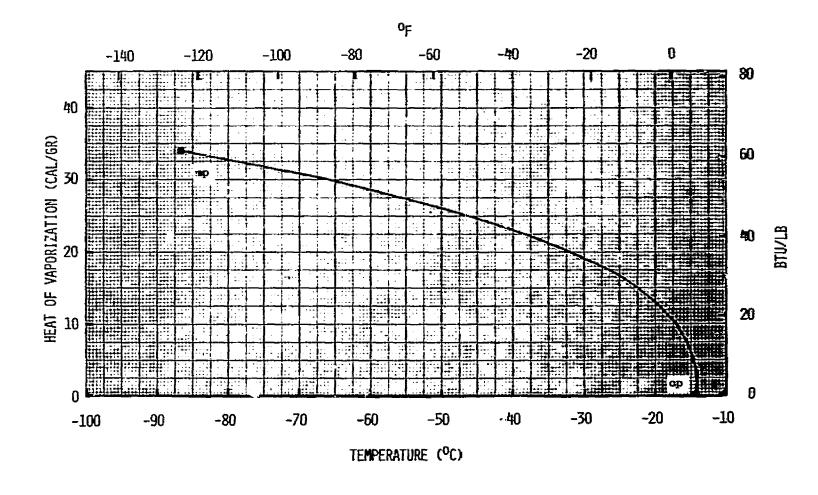


Figure 2.5-2 Heat of Vaporization vs Temperature for Silicon Tetrafluoride

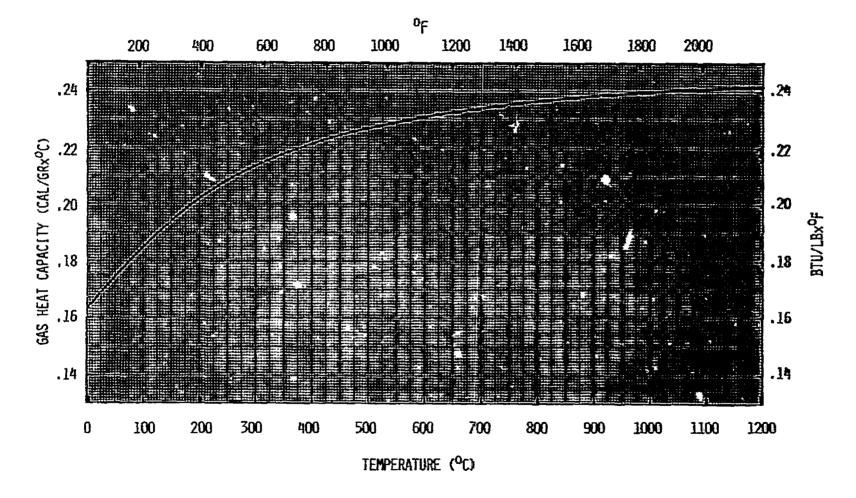


Figure 2.5-3 Gas Reat Capacity vs Temperature for Silicon Tetrafluoride

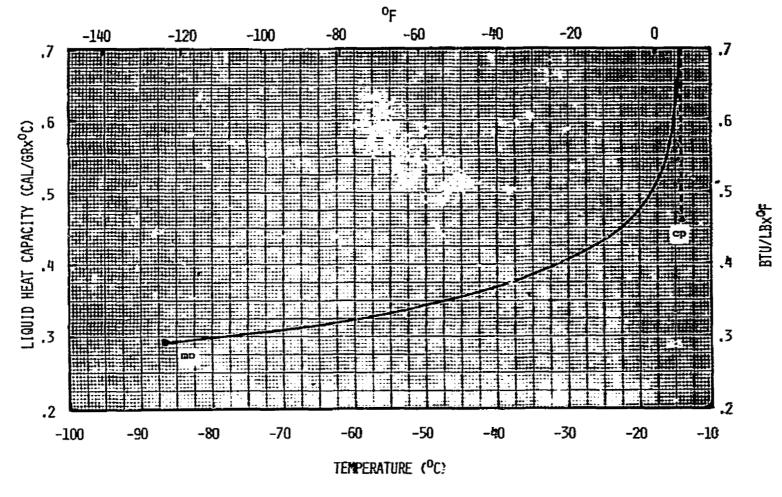


Figure 2.5-4 Liquid Heat Capacity vs Temperature for Silicon Tetrafluoride

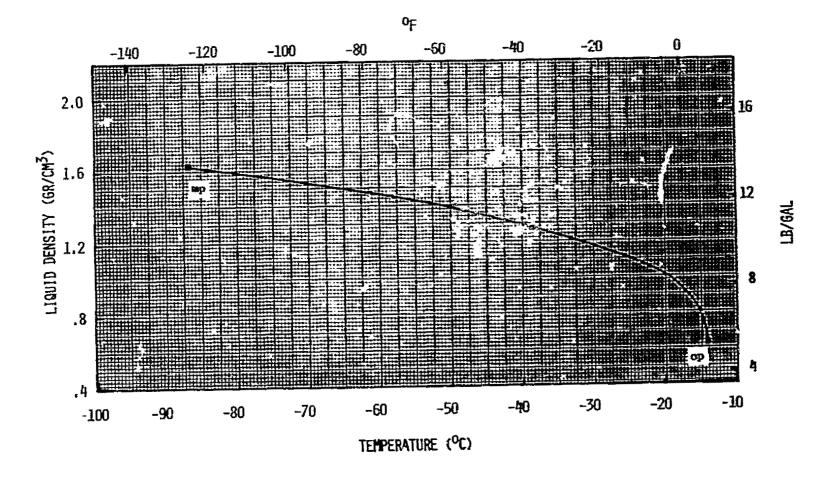


Figure 2.5-5 Liquid Density vs Temperature for Silicon Tetrafluoride

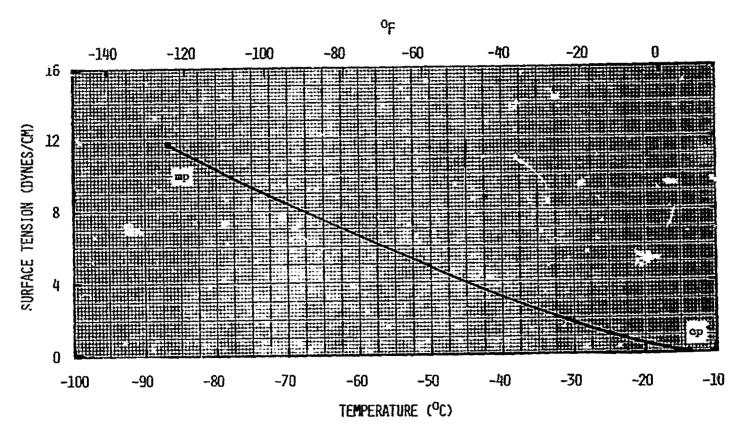


Figure 2.5-6 Surface Tension vs Temperature for Silicon Tetrafluoride

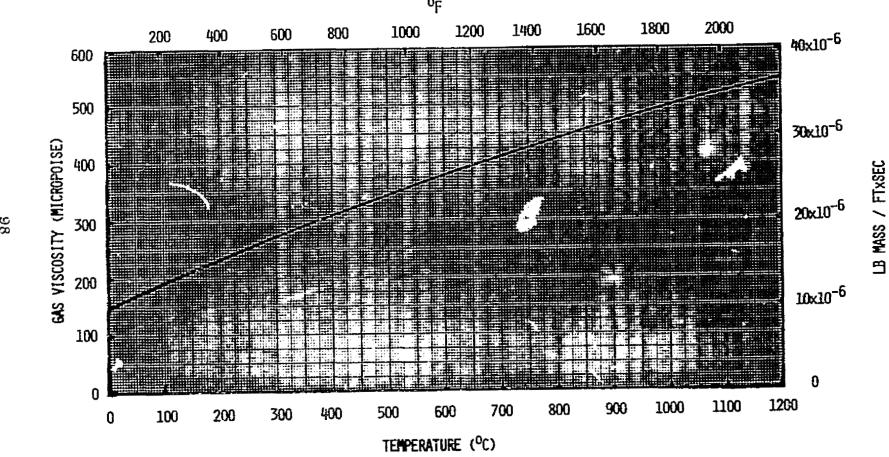


Figure 2.5-7 Gas Viscosity vs Temperature for Silicon Tetrafluoride

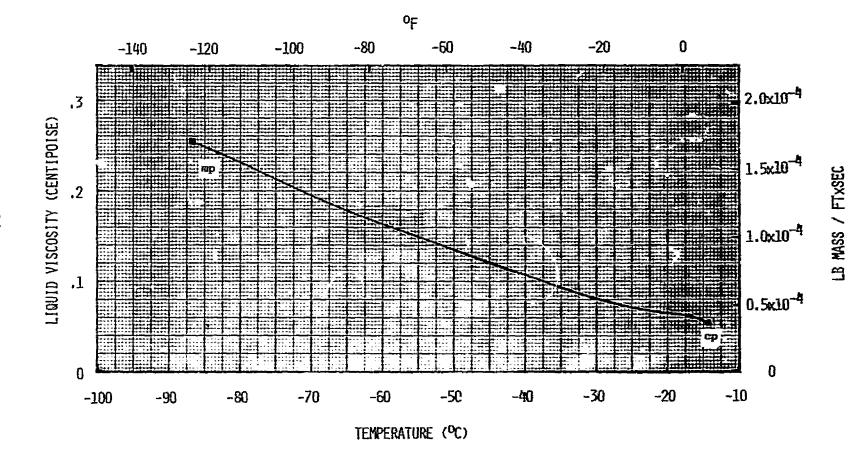
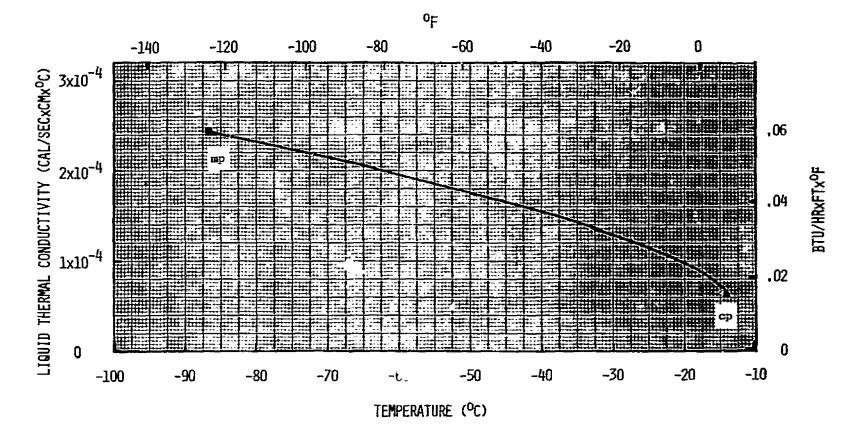


Figure 2.5-8 Liquid Viscosity vs Temperature for Silicon Tetrafluoride



Pigure 2.5-10 Liquid Thermal Conductivity vs Temperature for Silicon Tetrafluoride

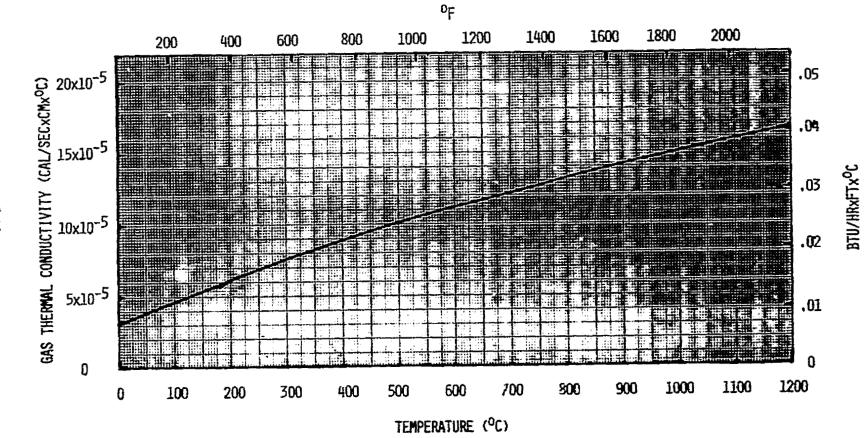


Figure 2.5-9 Gas Thermal Conductivity vs Temperature for Silicon Tetrafluoride

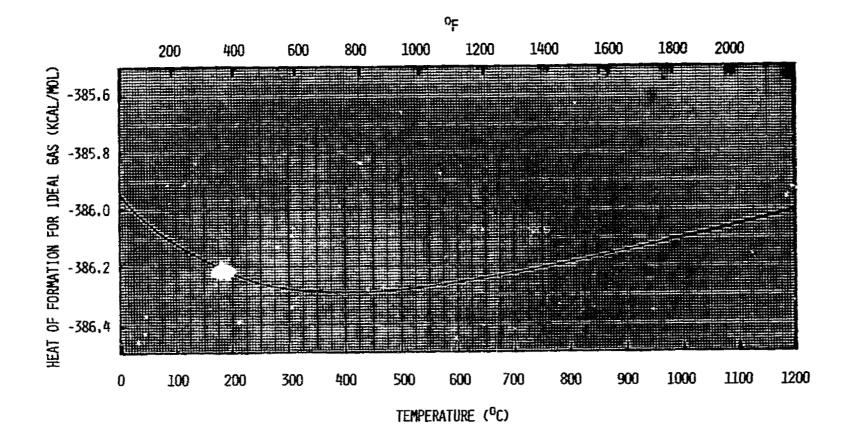


Figure 2.5-11 Heat of Formation vs Temperature for Silicon Tetrafluoride

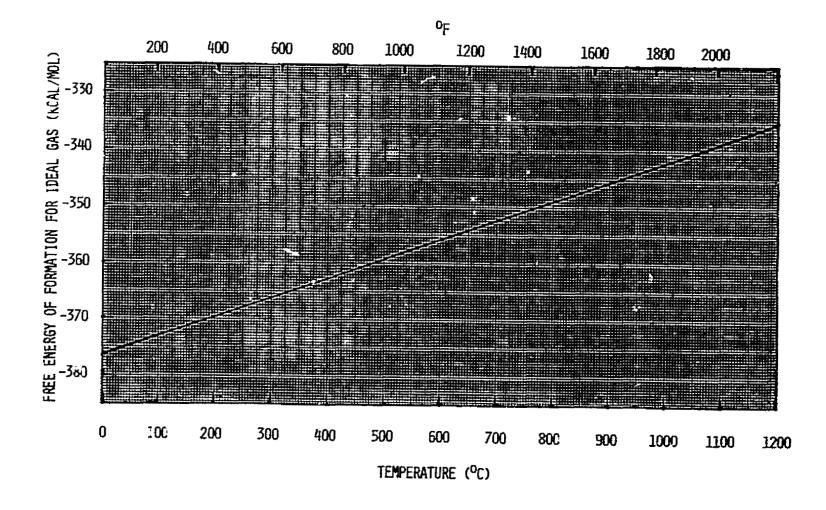


Figure 2.5-12 Free Energy of Formation vs Temperature for Silicon Tetrafluoride

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2.6 Silicon Properties

PHYSICAL PROPERTIES (TABLE 2.6-1)

Experimental values for the melting point have been reported (1, 32, 49, 67, 110); however, all other values have been calculated. Estimated values for the boiling point range from 2285°C to 3267°C (10, 31, 52, 55, 109, 124, 125, 137). Our value is estimated to give a reasonable computer fit to the available vapor pressure data. Estimated values of critical properties are reported by van Laar's calculation (5, 34, 43, 44, 101, 125), Baibus (4) and Gates and Thodos (31, reported in Table I-1). Solid properties listed in Table I-1 are at room temperature while liquid properties are at the melting point.

VAPOR PRESSURE (FIGURE 2.6-1)

Recent vapor pressure data reported by American (1, 18) and British (6) workers were selected and extended using the YSSP vapor pressure correlation (157):

$$\log P_V = A + \frac{B}{T} + C \log T + DT + ET^2$$
 (2.6-1)

where P_V is the vapor pressure of saturated liquid, mm Hg; T is temperature, *K; and A, B, C, D and E are correlation constants derived using a generalized least squares program. Other data (10, 125, 142) were not used because of high percentage error which is reported to be due to extensive reaction of the silicon (54, 165). For the 44 experimental data points used (which are all in the range below 0.2 mm Hg) the average absolute deviation was 17%.

HEAT OF VAPORIZATION (FIGURE 2.6-2)

Heat of vaporization values of about 3850 cal/gm are available (1, 6, 18, 55) as well as older (10, 125a, 147) and Russian (142) values of about 3170 cal/gm. From the vapor pressure data near the melting point, the heat of vaporization was determined using the Clausius-Clapyrin equation. Using these values, Watson's correlation (165) was used to extend the heat of vaporization to the boiling point.

$$\Lambda H_{V} = \Lambda H_{V_{1}} \left[\frac{7500 - T}{7500 - T_{1}} \right]^{n}$$
 (2.6-2)

where n=0.38 and all other terms have their usual meanings. The calculated values give a 1.3% absolute percentage deviation with the five experimental references giving values near 3850 cal/gm near the melting point.

HEAT OF SUBLIMATION (FIGURE 2.6-3)

Heat of sublimation based on limited data have been reported recently in the literature (18, 54, 106). Using the YSSP correlation of vapor pressure data (as described earlier), heats of sublimation were calculated using the Clausius-Clapeyron equation (123):

$$\Delta H_{\text{sub}} = P\Delta V_{\text{sub}} \frac{\delta P}{\delta T}$$
 (2.6-3)

where $\Delta H_{\rm sub}$ is the heat of sublimation, cal/gr-mol; P is the vapor pressure, atm; and $\Delta V_{\rm sub} = V_{\rm gas} - V_{\rm solid}$. The derivative, $\delta P/\delta T$, was determined from differentiation of the YSSP vapor pressure equation. Considering the possible inaccuracy in the extrapolation of very low vapor pressures at low temperatures, these values should be considered only order-of-magnitude calculations below 600°C.

HEAT CAPACITY (FIGURES 2.6-4 and 2.6-5)

Liquid heat capacities have been reported from experiments done in the range from the melting point to about 200°C above the melting point (67, 100). The values of Kantur (67) were selected because the temperature range was significantly greater with the temperatures appearing to be more accurately determined. The average values of heat capacity and temperature were taken as a reference point and the values were extended over the liquid range using the relationship:

Liquid Heat Capacity x Liquid Density = constant (2.6-4)

Calculated values agree within two percent of the values published in the experimental work (67).

Solid heat capacities have been reported by many authors (144, 138, 55, 34, 96, 164, 62, 115, and others) which give similar values. The JANAF and Touloukian values (138, 144) were selected.

DENSITY (FIGURES 2.6-6 & 2.6-7)

Several authors (93, 37, 14, 25, 9) have reported measured liquid density values from the melting point to about 400°C above the melting point. The datawere extended to the boiling point using a modification of the Rackett equation (166):

$$\rho = \rho_{\rm c} z^{-(1 - T_{\rm r})^{2/7}}$$
 (2.6-3)

where ρ_c is critical density; T_r is reduced temperature and z is a parameter defined by the experimental data. Calculated values give a 0.5% average absolute deviation from the thirty reported experimental values. The vertical line at the melting point indicates the change in density upon melting.

Solid density measurements of silicon are recorded near room temperature (133, 134) at the melting point (85) and many others give linear thermal expansion data which are summarized by Touloukian (144). Solid densities at various temperatures were calculated using the percentage linear expansions (144) according to the relation:

$$\rho = \rho_1 \times \{1 - [3x(percent linear expansion)]\}$$
 (2.6-6)

Calculated values gave less than one percent deviation with the measured values over the solid range.

SURFACE TENSION (FIGURE 2.6-8)

Limited data for the surface tension of silicon show a wide range of values (79, 71, 130, 25, 40). From the experimental data in close agreement (79, 71, 130), values were extended to the boiling point using the Othmer relation (123):

$$\sigma = \sigma_1 \left[\frac{7500 - T}{7500 - T_1} \right]^n \tag{2.6-7}$$

where σ is surface tension at T_1 , dynes/cm, and n is the correlation parameter, 1.2. The other parameters have their usual meaning. Calculated values agree with the limited data (5 values) with a 1.5% absolute error.

LIQUID VISCOSITY (FIGURE 2.6-9)

Liquid viscosity data for silicon are available from the melting point to about 400°C above the melting point (9, 81, 127, 163). Values from the melting point to the boiling point were calculated using a log η_L vs 1/T linear relationship. Average absolute percentage error was 9.1% on 19 data points, due largely to the wide scatter of the experimental data.

THERMAL CONDUCTIVITY (FIGURE 2.6-10)

An experimental value of the liquid thermal conductivity of silicon has been reported by Russian workers (167). Their research indicates a value of 0.16 (\pm .02) cal/cm \times sec \times °C at the melting point. The higher value of the thermal conductivity of liquid silicon compared to solid silicon at the melting point is in agreement with other experimental work (168).

Solid thermal conductivity data has been reported by several authors (104, 23, 83 and others). The recommended values of Touloukian (144) were selected.

TABLE 2.6-1

PHYSICAL PROPERTIES AND CRITICAL CONSTANTS OF SILICON

No.	<u>Identification</u>	<u>silicon</u>
1.	Symbol	Si
2.	State (std. cond.)	Solid
3.	Atomic Weight	28.086
4.	Boiling Point, b.p., °C	2,878*
5.	Melting Point, m.p., °C	1,412 ± 2
5,	Critical Temperature, To, °C	4,886*
7.	Critical Pressure, Pc, atm	530*
8.	Critical Volume, V _G , cm ³ /gr mol	232.6*
9.	Critical Density, pc, gr/cm3	0.1207*
10.	Vapor Pressure, mm Hg	2.8x10 ⁻⁴ (at m.p.)
11.	Heat of Vaporization, cal/gr	3,812 (at m.p.)
12.	Heat of Sublimation, cal/gr	4,075 (at m.p.)
13.	Heat of Fusion, cal/gr	264* (at m.p.)
14.	Liquid Heat Capacity, cal/gr-mol °C	6.755 (at m.p.)
15.	Solid Heat Capacity, cal/gr-mol °C	4.78 (at 25°C)
16.	Liquid Density, gr/cm ³	2.533 (at m.p.)
17.	Solid Density, gr/cm ³	2.329 (at 25°C)
18.	Percent Expansion on Freezing	10% (at m.p.)
19.	Surface Tension, dynes/cm	736 (at m.p.)
20.	Liquid Viscosity, centipoise	0.88 (at m.p.)
21.	Liquid Thermal Conductivity, cal/secxcmx°C	0.16 (at m.p.)
22.	Solid Thermal Conductivity, cal/secxcmx°C	0.353 (at 25°C)

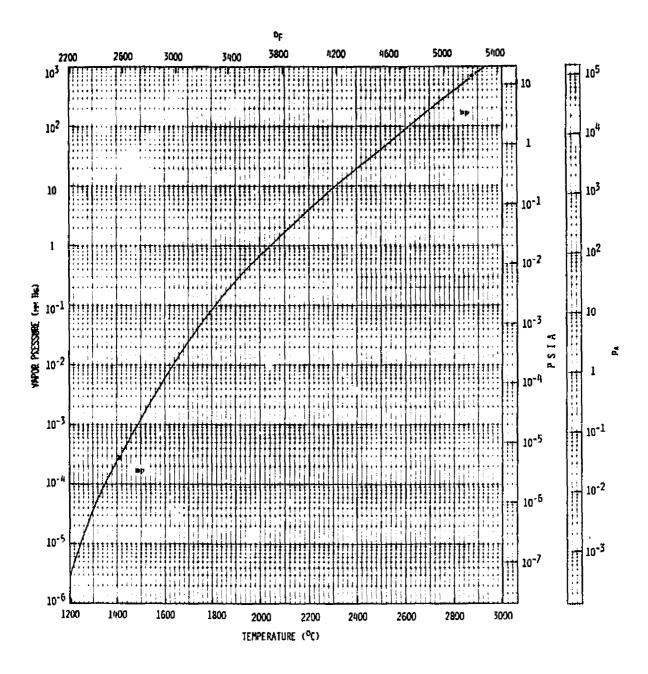


Figure 2.6-1 Vapor Pressure vs Temperature for Silicon

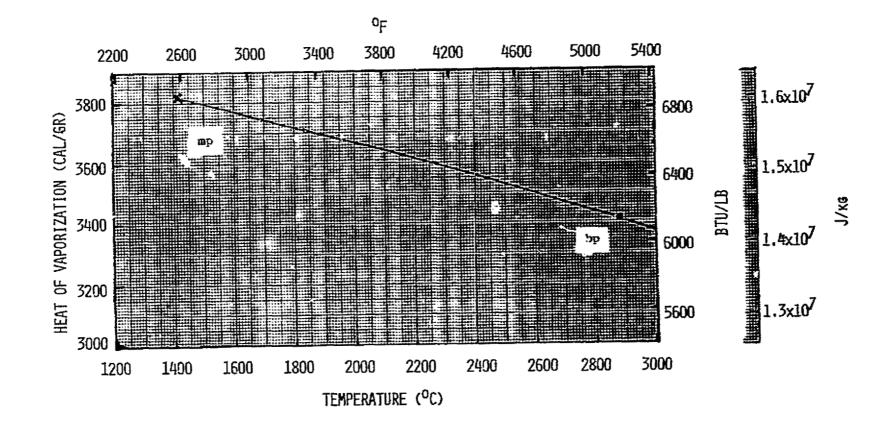


Figure 2.6-2 Heat of Vaporization vs Temperature for Silicon

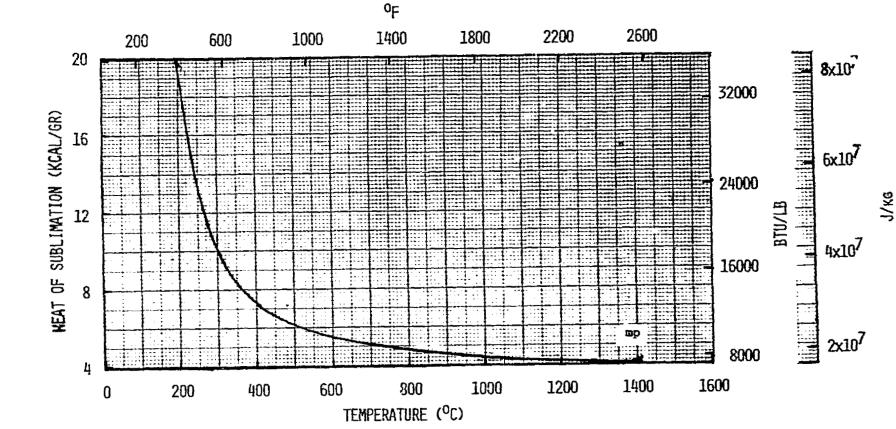


Figure 2.6-3 Heat of Sublimation vs Temperature for Silicon

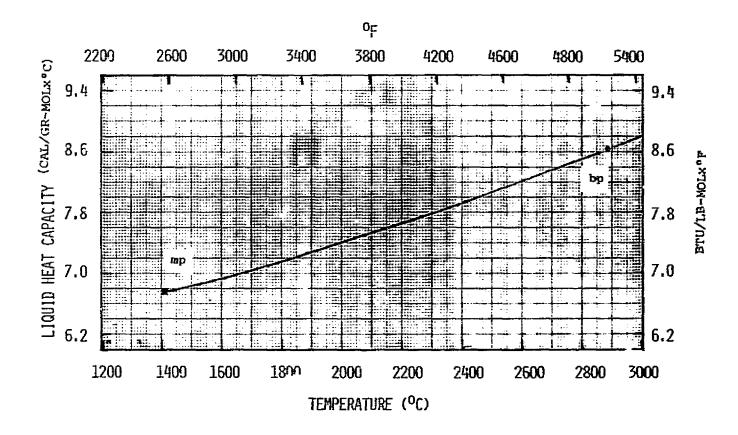


Figure 2.6-4 Liquid Heat Capacity vs Temperature for Silicon

Figure 2.6-5 Solid Heat Capacity vs Temperature for Silicon

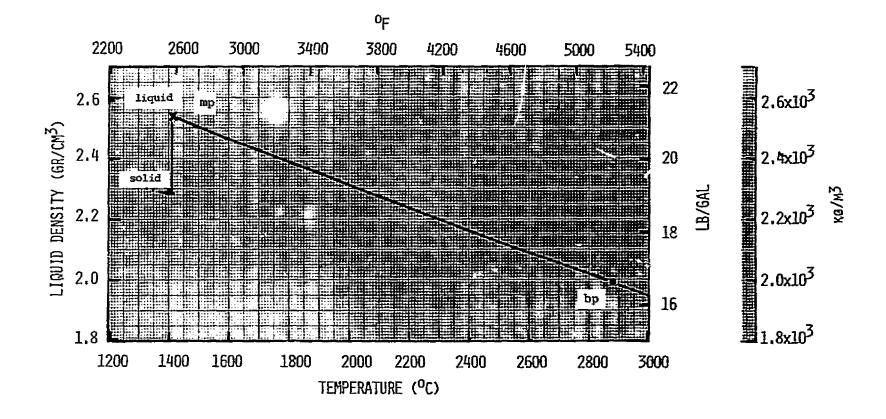


Figure 2.6-6 Liquid Density vs Temperature for Silicon

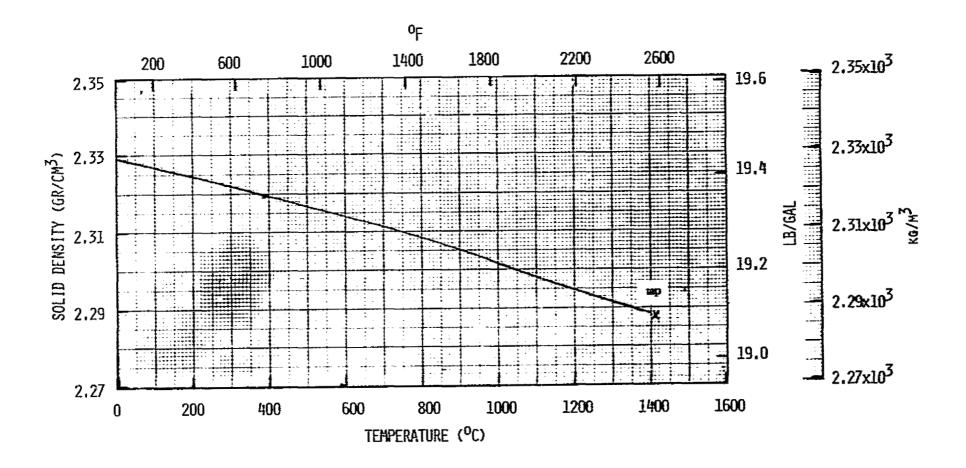


Figure 2.6-7 Solid Density vs Temperature for Silicon

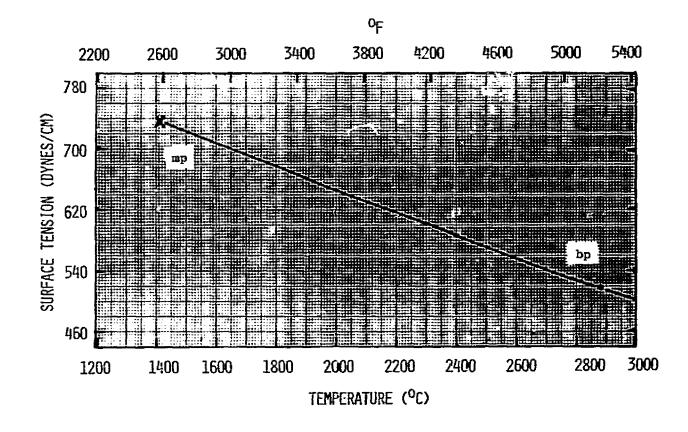


Figure 2.6-8 Surface Tension vs Temperature for Silicon

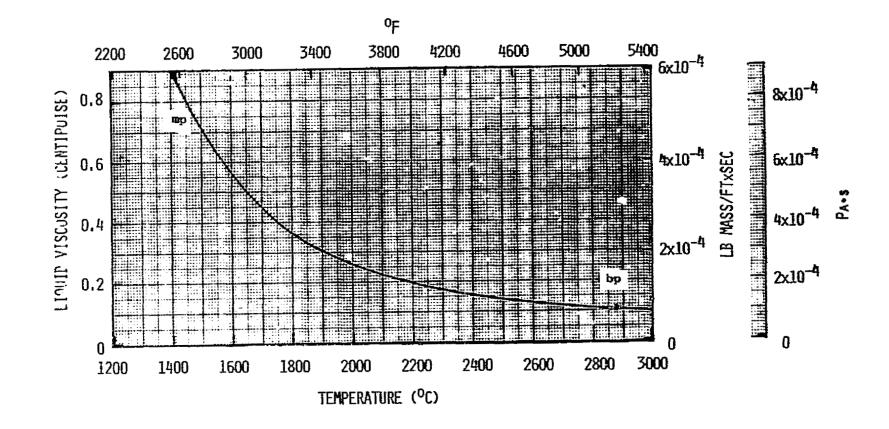


Figure 2.6-9 Liquid Viscosity vs Temperature for Silicon

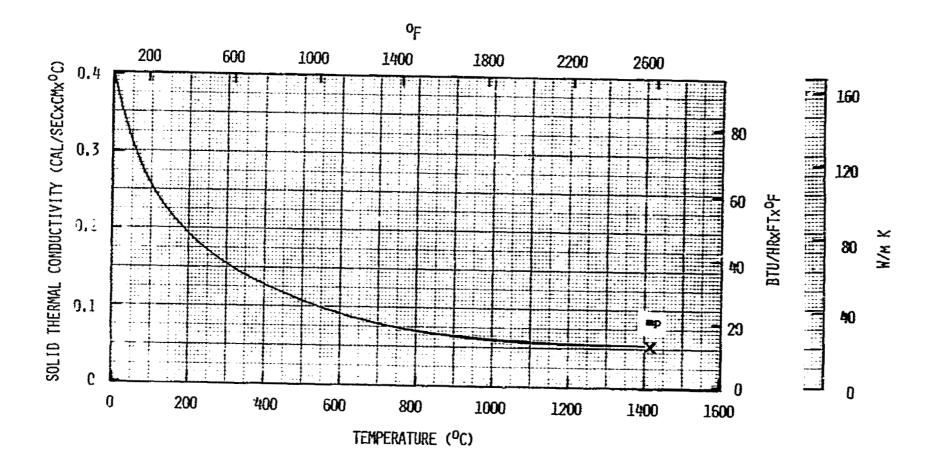


Figure 2.6-10 Solid Thermal Conductivity vs Temperature for Silicon

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2.7 Thermal Conductivity Investigation

Gas phase thermal conductivity values were experimentally determined between 25°C and 350°C for a variety of silicon source materials which included silane and halogenated solanes. The apparatus used was a hot wire thermal conductivity cell (or catharometer). It consists of two pairs of matched tungsten-rhenium filaments mounted in a stainless steel block. The filaments are connected as elements of a constant current Wheatstone Bridge (Figure 2.7-1). The cell is electrically heated and a constant temperature is maintained with a digital temperature controller and read-out to + 1°C. The filaments are positioned in cavities in the steel block into which the gases, of which the thermal conductivity is to be determined, can be introduced. The filaments are heated by a constant current and the heat thus generated is dissipated primarily by conduction through the gas. A change in the thermal conductivity of the gaseous medium results in a change in the rate of dissipation and therefore, a change in the temperature of the filament. The temperature of the hot filament is measured as if it were a resistance thermometer; change in temperature produces a change in filament resistance, which is measured by means of the Wheatstone Bridge circuit.

The thermocouples (type K) used to monitor the temperature of the thermal conductivity cell were calibrated using materials of known melting points throughout the temperature range of the study (25°C to 350°C). The EMF of the thermocouples was measured with a Leeds and Northrup, Model 8686, millivolt potentiometer which was calibrated and certified at the factory. The temperatures reported for the thermal conductivity values are considered to be accurate to +1°C.

Since absolute measurement of thermal conductivity is difficult, a differential method was employed in which the catharometer was divided into two parts where half of the filaments are in contact with a reference gas of known thermal conductivity and the other half contact the sample whose thermal conductivity is to be determined. The Wheatstone Bridge is first balanced by introducing the reference gas into both sides of the cell. The sample to be determined is then introduced into the sample side of the cell and the resultant voltage unbalance (E) is recorded. The catharometer responds to the reciprocal of the thermal conductivities according to equation 2.7-1: (reference 1)

$$E - E_{ref.} = b(^{1}/\lambda - ^{1}/\lambda_{ref}) \qquad (2.7-1)$$

where $E_{\rm ref}$ is voltage with the reference gas in both sides of the thermal conductivity cell, λ and $\lambda_{\rm ref}$ are the thermal conductivities of the unknown and reference gas respectively, and by a constant characteristic of the paricular apparatus (cell constant). This cell constant (b) can be determined by using a standardization gas of known thermal conductivity

as the sample and determining the voltage unbalance (E) of it with respect to the reference gas. The cell constant (b) is slightly temperature dependent and must be determined throughout the temperature range in which measurements are to be made.

Before thermal conductivity data could be obtained, the apparatus described above needed to be calibrated. The calibration work included the determination of cell constants for the temperature range 25°C to 350°C, the determination of filament wire temperatures for various filament currents and cell wall temperatures, and the experimental determination of the thermal conductivity of argon and hydrogen in the temperature range 25°C to 350°C.

The ceil constant, which is used to calculate thermal conductivity values when the differential method is used, is temperature dependent and therefore needs to be determined for the complete temperature range to be investigated. It was also found that at a given temperature, the cell constant may vary slightly from day to day; therefore cell constants were routinely determined everytime data were collected. This variation may be due to slight changes in the filament current or to slight oxidation or corrosion of the filament with use.

In measuring the thermal conductivity of gases using the "hot wire" method, the gas may not be at a uniform temperature due to differences in the temperature of the cell wall and filament wire. This can be minimized by operating the apparatus at filament currents sufficiently low that this temperature difference is small. In order to do this, a means of monitoring the filament wire temperature was needed. was accomplished by using the filament as a resistance thermometer. With no current in the filament, the filament resistance as a function of temperature was measured (figure 2.7-2). When thermal conductivity data were being obtained, the filament resistance was routinely calculated by monitoring the current through the filament and the potential across the filament. The filament temperature can then be obtained from figure 2.7-1. The filament current was then adjusted so that the temperature difference between the filament and the cell wall was small.

The thermal conductivity of argon was determined through out the temperature range 25°C to 350°C. These values were compared to recommended values for the thermal conductivity of argon (reference 2) in order to evaluate the accuracy of data obtained on this apparatus (figure 2.7-3). The recommended values used were those presented in "Thermophysical Properties of Matter", Vol. 3 on Thermal Conductivity (TPRC), and were determined by an evaluation of available published data. It was stated that the published data correlated with the recommended values to within \pm 5%. The thermal conductivity values obtained in this study agree with the recommended values to within \pm 4% from 300°C to 350°C.

The thermal conductivity of hydrogen was determined in the temperature range 25°C to 350°C. These values were compared to previously reported experimental values for thermal conductivity of hydrogen (references 3 and 4) in order to evaluate the accuracy of the data obtained on this apparatus (figure 2.7-4) for gases of relatively high thermal conductivity.

The thermal conductivity of silane (SiH₄) was determined between 25°C and 300°C (Table 2.7-1 and Figure 2.7-5). Values above 300°C were not determined because above that temperature silane is thermally unstable and begins to deposit silicon. There have been no previously reported experimental data for gaseous thermal conductivity of silane. Estimated values have been determined using a modified Eucken Correlation (ref. 5) and these estimated values agree fairly well with the now available experimental values.

The thermal conductivity of dichlorosilane (SiH₂Cl₂) was determined between 25°C and 350°C (Table 2.7-2 and Figure 2.7-6). There have been no previously reported experimental data for gaseous thermal conductivity of dichlorosilane. Estimated values have been reported (reference 39) in the temperature range 0°C to 70°C which were determined by a Euchen approximation. These estimated values are considerably lower than the experimental values now reported.

The thermal conductivity of trichlorosilane (SiHCl₃) was determined between 50°C and 350°C (Table 2.7-3 and Figure 2.7-7). There have been no previously reported experimental data for gaseous thermal conductivity of trichlorosilane.

The thermal conductivity of gaseous silicon tetrachloride was determined between 100°C and 350°C (Table 2.7-4 and Figure 2.7-8). There have been both calculated (ref. 7) and experimental (ref. 8) values for the thermal conductivity of silicon tetrachloride previously reported. The calculated values, in the temperature range 80°C to 335°C, were lower than the values obtained in this study by more than 10°C. The experimental values, in the temperature range 70°C to 300°C, were about 10°C lower than the values obtained in this study.

The thermal conductivity of silicon tetrafluoride (SiF_A) was determined beween 25° and 350°C (Table 2.7-5 and Figure 2.7-9). The values obtained in this study agree to within $\pm 3\%$ with previously reported (reference 9) experimental data for silicon tetrafluoride (figure 2.7-10)

Figure 2.7-11 summarizes all of the experimentally determined values for gaseous—thermal conductivity of silane and halogenated silanes reported from this investigation.

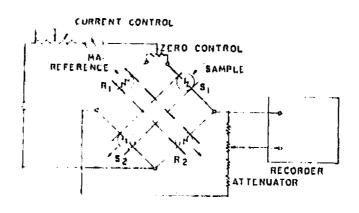


Figure 2.7-1 Wheatstone Bridge Circuit For Thermal Conductivity Cell

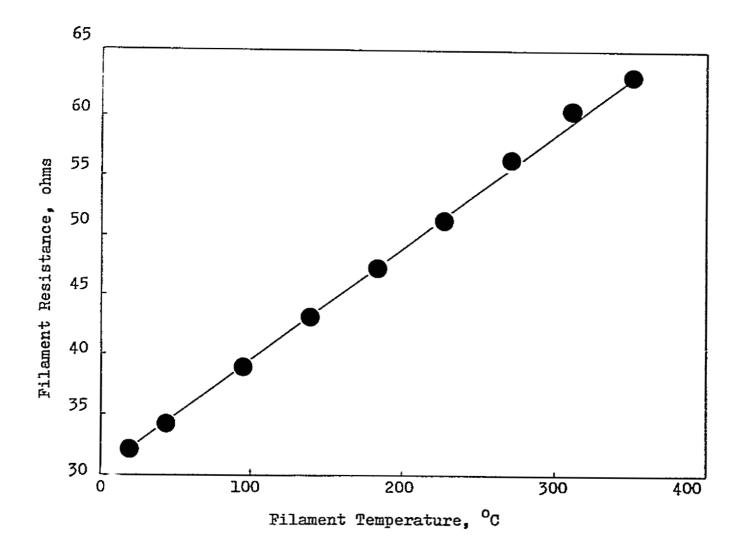


Figure 2.7-2 Filament Resistance as a Function of Temperature

Figure 2.7-3 Comparison of Thermal Conductivity Values for Argon

Figure 2.7-4 Comparison of Thermal Conductivity Values for Hydrogen

Table 2.7-1 Gaseous Thermal Conductivity Values of Silane

Temperature	Gaseous Thermal Conductivity		
<u>°c</u>	mW cm-l oK-l	Cal cm ^{-l} sec ^{-l o} C ^{-l}	BTU hr-lft-l og-l
28.0	0.234	56.02 X 10 ⁻⁶	13.54 X 10 ⁻³
45.7	0.249	59.44 X 10 ⁻⁶	14.37 X 10 ⁻³
94.7	0.297	70.96 X 10 ⁻⁶	17.15 X 10 ⁻³
139.4	0.345	82.34 X 10 ⁻⁶	19.90 X 10 ⁻³
184.1	0.400	95.67 x 10 ⁻⁶	23.13 X 10 ⁻³
227.4	0.449	107.24 x 10 ⁻⁶	25.93 X 10 ⁻³
269.5	0.497	118.86 X 10 ⁻⁶	28.73 X 10 ⁻³

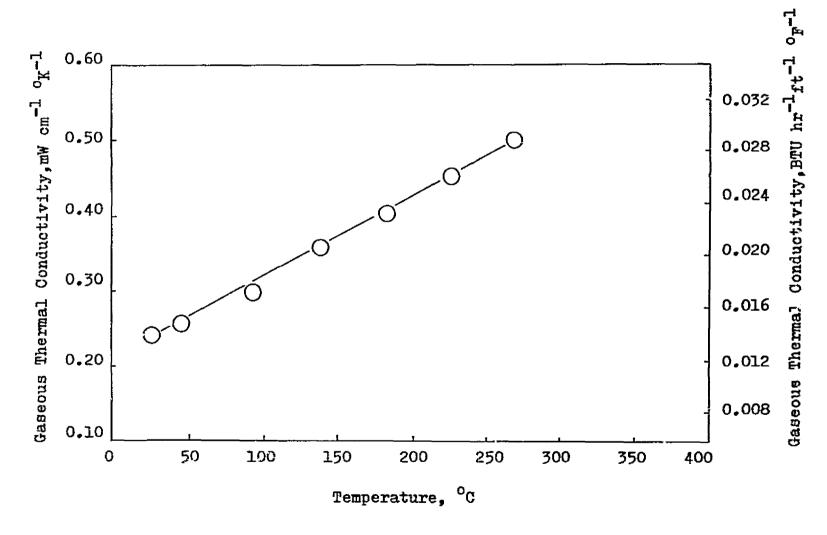


Figure 2.7-5 Gaseous Thermal Conductivity of Silane

Table 2.7-2 Gaseous Thermal Conductivity Values of Dichlorosilane

<u>Temperature</u>	Gaseous Thermal Conductivity		
<u>°c</u>	mw cm-l oK-l	Cal cm-lsec-l oc-l	BTU hr-1ft-1 op-1
28.0	0.102	24.43 X 10 ⁻⁶	5.91 X 10 ⁻³
45.7	0.108	25.72 X 10 ⁻⁶	6.22×10^{-3}
94.7	0.129	30.86 X 10 ⁻⁶	7.46 \times 10 ⁻³
139.4	0.148	35.42 X 10 ⁻⁶	8.56 X 10 ⁻³
184.1	0.169	40.37 × 10 ⁻⁶	9.76×10^{-3}
2.7.4	0.194	46.46×10^{-6}	11.23 X 10 ⁻³
9.5	0.217	51.79 X 10 ⁻⁶	12.52 X 10 ⁻³
11.3	0.243	58.15 X 10 ⁻⁶	14.06×10^{-3}
.50.6	0.267	63.70 x 10 ⁻⁶	15.40 X 10 ⁻³

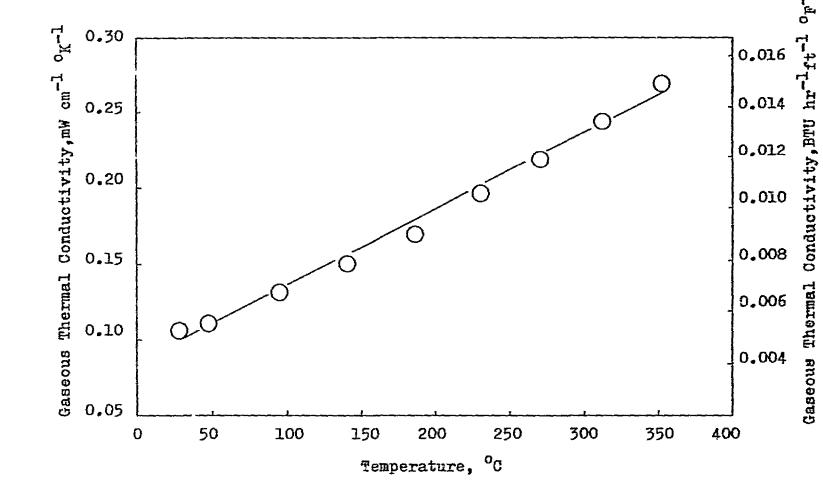


Figure 2.7-6 Gaseous Thermal Conductivity of Dichlorosilane

Table $2 \cdot 7 - 3$ Gaseous Thermal Conductivity Values of Trichlorosilane

	Temperature		aseous Thermal Conducti	vity
	<u>°c</u>	mW cm ⁻¹ o _K -1	Cal cm ^{-l} sec ^{-l c} c ^{-l}	BTU hr-lft-l cF-l
	45.7	0.093	22.13 X 10 ⁻⁶	5.35 X 10 ⁻³
142	94.7	0,110	26.22 X 10 ⁻⁶	6.34×10^{-3}
	139.4	0.126	30.16 X 10 ⁻⁶	7.29 X 10 ⁻³
	184.1	0.144	34.35 X 10 ⁻⁶	8.30 X 10 ⁻³
	227.4	0.161	38.55 x 10 ⁻⁶	9.32 X 10 ⁻³
	269.5	0.180	43.05 x 10 ⁻⁶	10.41×10^{-3}
	311.3	0.198	47.24 X 10 ⁻⁶	11.42 X 10 ⁻³
	350.6	0.216	51.58 X 10 ⁻⁶	12.47×10^{-3}

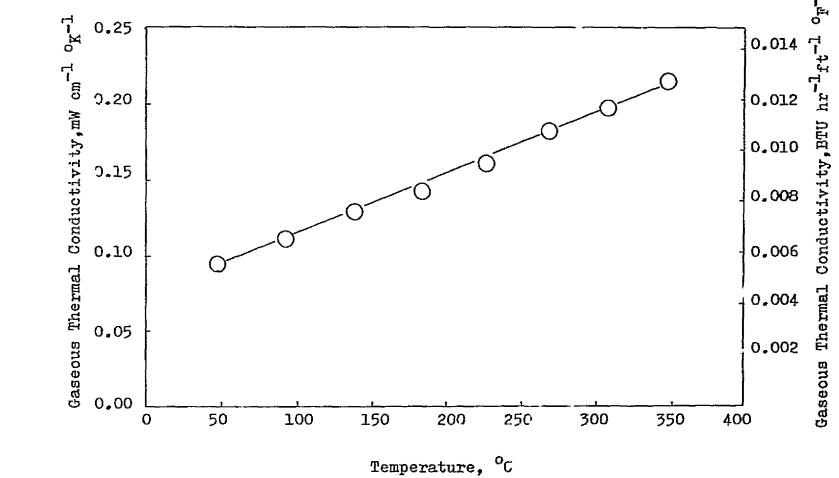


Figure 2.7-7 Gaseous Thermal Conductivity of Trichlorosilane

Table 2.7-4 Gaseous Thermal Conductivity Values of Tetrachlorosilane

Temperature	Gaseous Thermal Conductivity		
<u>°с</u>	$mW cm^{-1} oK^{-1}$	Cal cm-lsec-l oc-l	BTU hr-1ft-1 oF-1
94.7	0.100	23.93 × 10 ⁻⁶	5.78 X 10 ⁻³
139.4	0.111	26.43×10^{-6}	6.39 X 10 ⁻³
184.1	0.124	29.59 X 10 ⁻⁶	7.15 \times 10 ⁻³
227.4	0.138	32.89 X 10 ⁻⁶	7.95 X 10 ⁻³
269.5	0.153	36.59 X 10 ⁻⁶	8.85 X 10 ⁻³
311.3	0.169	40.39 X 10 ⁻⁶	9.76 x 10 ⁻³
350.6	0.193	46.13 X 10 ⁻⁶	11.15 X 10 ⁻³

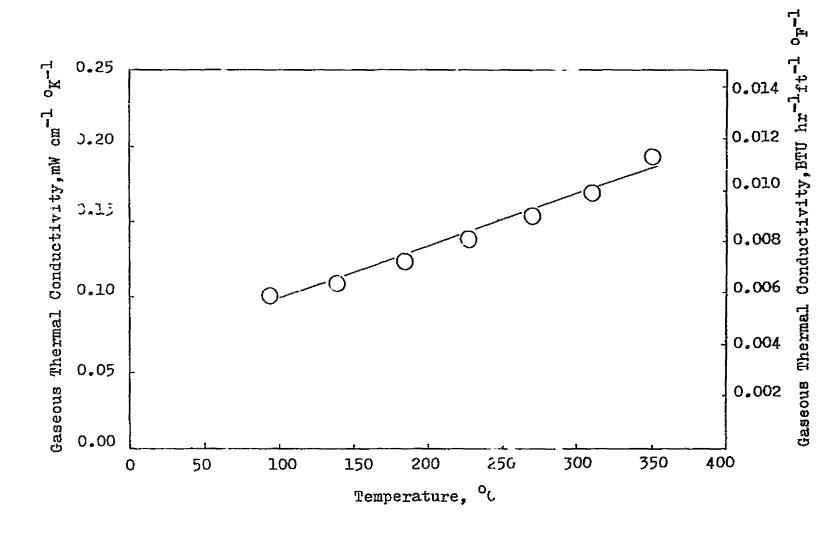


Figure 2.7-8 Gaseous Thermal Conductivity of Tetrachlorosilane

Table 2.7-5 Gaseous Thermal Conductivity Values of Tetrafluorosilane

	<u>Temperature</u>	Gaseous Thermal Conductivity		
	<u>°c</u>	mW cm-l oK-l	Cal cm-lsec-l oc-l	BTU hr-lft-l og-l
	29.0	0.150	35.95 X 10 ⁻⁶	8.69 X 10 ⁻³
1-16	45.7	0.158	37.79 X 10 ⁻⁶	9.13 X 10 ⁻³
. ,	94.7	0.189	45.24 X 10 ⁻⁶	10.94 X 10 ⁻³
	139.4	0.215	51.43 X 10 ⁻⁶	12.43×10^{-3}
	184.1	0.241	57.67 X 10 ⁻⁶	13.94 X 10 ⁻³
	227.4	0.274	65.46 X 10 ⁻⁶	15.83 X 10 ⁻³
	269.5	0.291	69.55 x 10 ⁻⁶	16.81 X 10 ⁻³
	311.3	0.316	75.55 X 10 ⁻⁶	18.26 X 10 ⁻³
	350.6	0.345	82.34 X 10 ⁻⁶	19.90 X 10 ⁻³

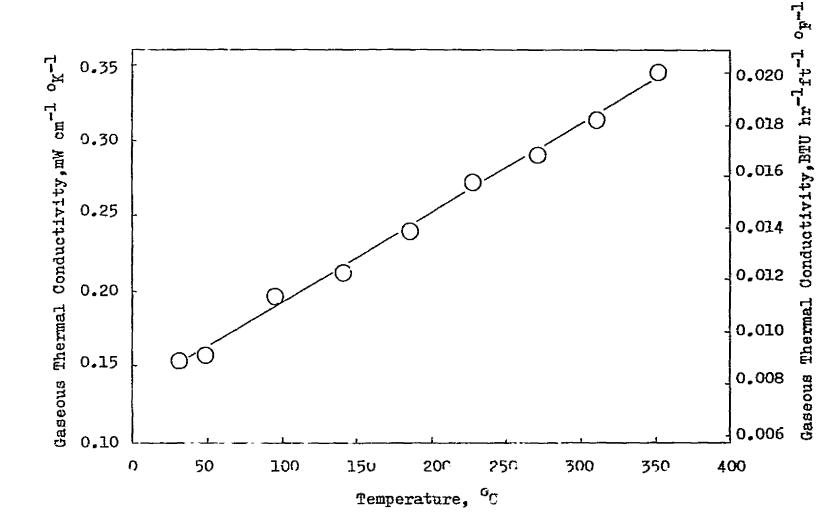


Figure 2.7-9 Gaseous Thermal Conductivity of Tetrafluorosilane

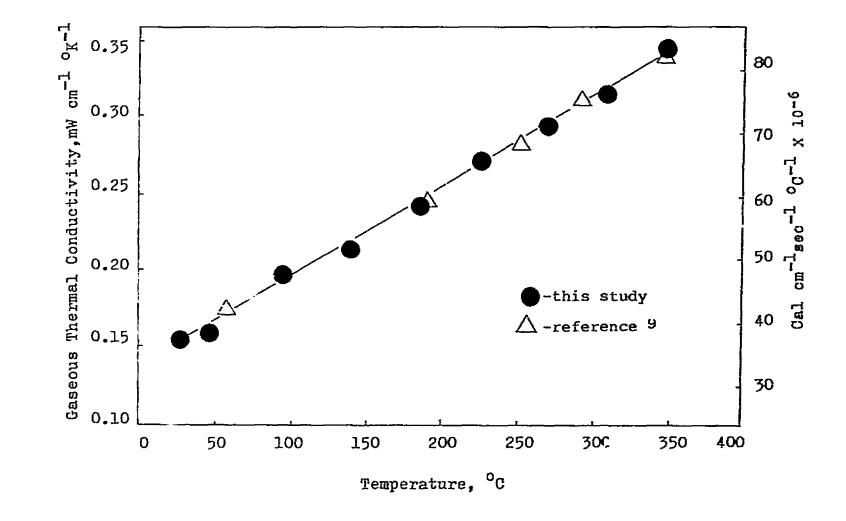


Figure 2.7-10 Comparison of Thermal Conductivity Values for Tetrafluorosilane

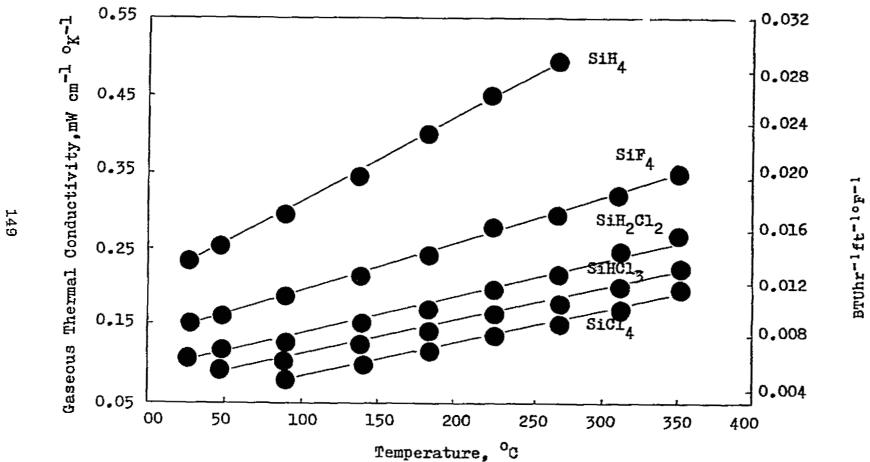


Figure 2.7-11 Gaseous Thermal Conductivity Values for Silane and Halogenated Silanes

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2.8 Viscosity Investigation

Gas viscosity values of some halogenated silanes were experimentally determined between 40°C and 200°C. The viscosity values were determined by a transpiration method which is based on the rate of flow of the gas through a capillary. In order to determine these gas flow rates, a constant volume, glass viscometer (Figure 2.8-1) was fabricated and assembled. The apparatus is similar to one described by McCoubrey and Singh (1). The apparatus consist of a 1 liter glass bulb which is connected through a glass manifold to a mercury manometer and to a thermostated capillary with a preheater coil. The capillary is 20 cm. in length and has an internal diameter of 0.02 cm. The exit side of the capillary can be continuously exacted by a two stage mechanical pump.

In order to make a measurement, the viscometer is thoroughly evacuated and then the gas sample is introduced into the bulb up to a pressure of about 18 cm. Hg. The gas sample is then evacuated through the capillary and the resulting rate of flew is monitored by recording the pressure decrease in the bulb with time. The pressure is measured with a standard f tube mercury manometer to 10.5mm. Hg.

The rate of flow of a gas through a capillary is dependent upon the coefficient of viscosity (n) of the gas. By combining Poiseuille's equation for laminar flow of a gas through a tabe and the ideal gas law equation: the relationship between pressure, time, and viscosity of a gas can be derived. Poiseuille's equation (2) for laminar gas flow is:

seuille's equation (2) for laminar gas flow is:
$$\frac{dV}{dt} = \frac{{}^{n}(P_{1}^{2} - P_{2}^{2})r^{4}}{16InP_{0}}$$
(2.8-1)

where dV/dt is the volume rate of gas flowing through the capillary, P_1 is the pressure at the capillary inlet, P_0 is the pressure at the capillary outlet, r is the radius of the capillary, L is the length of the capillary, and P_0 is the pressure at which the gas volume is measured. In this method where the gas is continuously evacuated with a pump, P_0 is negligible compared to P_1 and equation 2.8-1 reduces (8)

$$\frac{dV}{dt} = \frac{\pi P_1^2 r^4}{100 \pi P_0}$$
 (2.8-2)

From the ideal gas law, aV, the volume of gas at P_0 , passing through the capillary in unit time can be expressed in terms of dN, the number of acts of gas flowing through the capillary in unit time (equation 2.8-3).

$$dV = dN(RT/P_0)$$
 (2.8-3)

Substituting equation 2.8-3 into equation 2.8-2 gives equation 2.8-4:

$$\frac{dN}{dt} = \frac{{}^{9}P_{1}^{2}r^{4}}{16LnRT}$$
 (2.8-4)

As the gas is evacuated in a constant volume viscometer, the pressure decreases. Again using the ideal gas law,

$$dN = -dP_1(V/RT) \qquad (2.8-5)$$

where V is the volume being evacuated. Substituting equation 2.8-5 into equation 2.8-4 gives equation 2.8-6

$$dP_1 = \frac{-\pi P_1}{16U_0V} \frac{2r^4}{dt}$$
 (2.8-6)

The assumptions of the derivations are: constant volume of the system, ideal behavior of the gas, and laminar flow through the callibry. Since the pressure is measured with a U-tube manemeter the volume of the system will change by 10 to 20 ml. during an experiment. However, with a total volume of the viscometer of over 1 liter, this change can be neglected. The gases to be measured do not exhibit ideal behavior, but at pressures of less than 1 atmosphere their deviations should not be large. Laminar flow assumes zero velocity at the wall. A correction may need to be made for slip at the warl.

From equation 2.8-6, the viscosity (n) of a gas can be calculated from the slope of the line obtained by plotting $1/P_1$ versus to the applicant dimensions and the volume of the system a gas of known viscosity can be used to determine an apparatus constant which includes all the constant terms in equation 2.8-6. Alternatively, the calculation constant can be omitted and the viscosity of the unknown, relative to that of a reference gas, can be computed from the inverse ration of the slopes of the 1/P versus t graphs (equation 2.8-7).

$$\frac{\eta}{\eta_{\text{rof}}} = \frac{\text{slope}}{\text{slope}} \tag{2.8-7}$$

Evaluation and calibration of the gas viscometer was accomplished before data collection begun. Using argon as a

reference to determine a viscometer constant, experimental values for gas viscosity of nitrogen have been determined between 40°C and 200°C. These values were compared to recommended values for the gas viscosity of nitrogen (3) in order to evaluate the accuracy of data obtained on this viscometer (figure 2.8-2). The recommended values used were those presented in "Thermophysical Properties of Matter", Vol. 11 on viscosity (TPRC), and were determined by an evaluation of available published data. It was stated that the published data correlated with the recommended values to within ±2%. The viscosity values obtained for gaseous nitrogen in this study deviate from the recommended values by less than 2% from 40°C 200°C (figure 2.8-2).

The viscosity of trichlorosilane (SiHCl₃) has been determined between 40°C and 200°C (table 2.8-1 and figure 2.8-3). There have been no previously reported experimental values for gar viscosity of trichlorosilane in the temperature range of the study. Values at 0°C and 31°C were reported by Tel'chuk and Tubyanskuya (ref. 5).

The viscosity of dichlorosilane (SiH₂Cl₂) has been determined between 40°C and 200°C (table 2.8-2 and figure 2.8-4). The sample of dichlorosilane used for the measurements was semiconductor grade obtained from Union Carbide Corporation. There have been no previously reported experimental values for the gas phase viscosity of dichlorosilane. One set of calculated values have been reported (ref. 6) in the temporature range of 0°C to 300°C. These calculated values agree with the experimental values determined in this study with deviations of less than ±2% from 40°C to 200°C.

The viscosity of tetrafluorosilane (SiF₄) has been determined between 40°C and 200°C (Table 2.8-3 and Figure 2.8-5). There have been two previous reports of experimentally determined viscosity values for tetrafluorosilane. Ellis and Raw (reference 4) reported values between 23°C and 134°C and Mc-Coubrey and Singh (reference 1) reported values between 18°C and 190°C. The values of McCoubrey and Singh were in close agreement to the values reported in this study with less than 3% deviation through the whole temperature range. The values of Ellis and Raw were lower than the values reported in this study by as much as 7%.

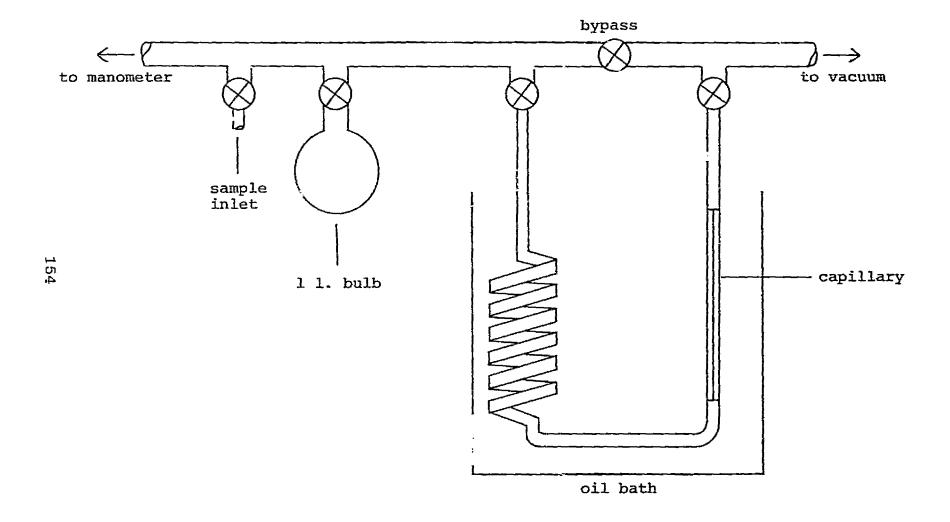
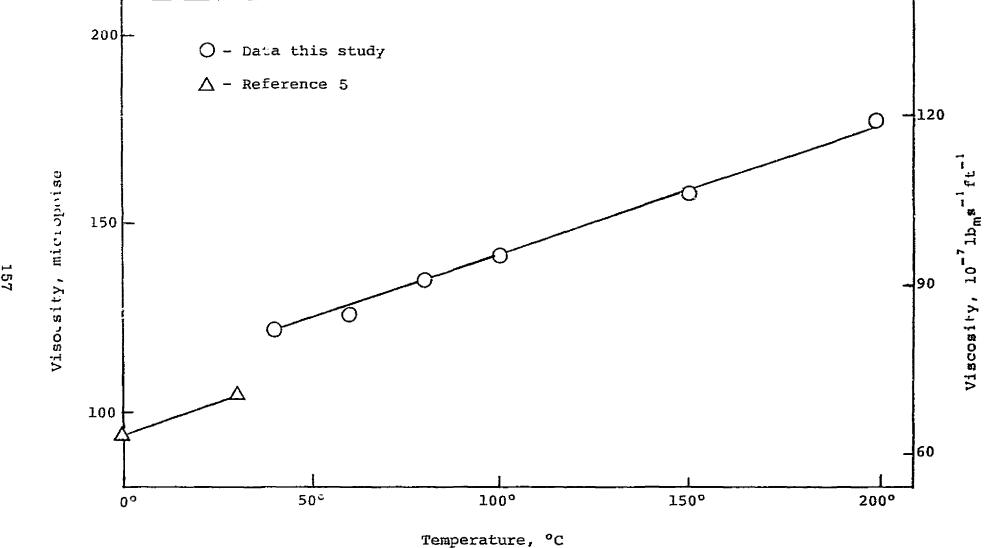


Figure 2.8-1 Constant Volume Gas Viscometer

Temperature, °C

Figure 2.8-2 Viscosity of Nitrogen

Temperature		Viscosity		
°c	micropoise	Nsm ⁻¹	lb s ⁻¹ ft ⁻¹	
÷ 0	122.0	12.20x10 ⁻⁵	8.20x10 ⁻⁶	
60	125.9	12.59x10 ⁻⁶	8-46x10 ⁻⁶	
80	134.8	13.48x10 ⁻⁶	9-06x10 ⁻⁶	
100	140.8	1'.08x10 ⁻⁶	9.46x10 ⁻⁶	
150	157.3	15.73x10 ⁻⁶	10.57x10 ⁻⁶	
200	177.2	17.72x10 ⁻⁶	11.91x10 ⁻⁶	



2.8-3 Viscosity of Gaseous Trichlorosilane Figure

Table 2.8-2 Viscosity of Gaseous Dichlorosilane

Temperature		Viscosity		
	³ C	micropolas	Nsm ⁻¹	lb _m s ⁻¹ ft ⁻¹
	40	118.7	11.87 x 10 ⁻⁶	7.98 X 10 ⁻⁶
158	60	125.3	12.53×10^{-6}	8.42×10^{-6}
	80	134.6	13.46 x 10 ⁻⁶	9.05×10^{-6}
	100	140.2	14.02×10^{-6}	9.42×10^{-6}
	150	163.5	16.35 x 10 ⁻⁶	10.99 X 10 ⁻⁶
	200	181.9	18.19 x 10 ⁻⁶	12.22×10^{-6}

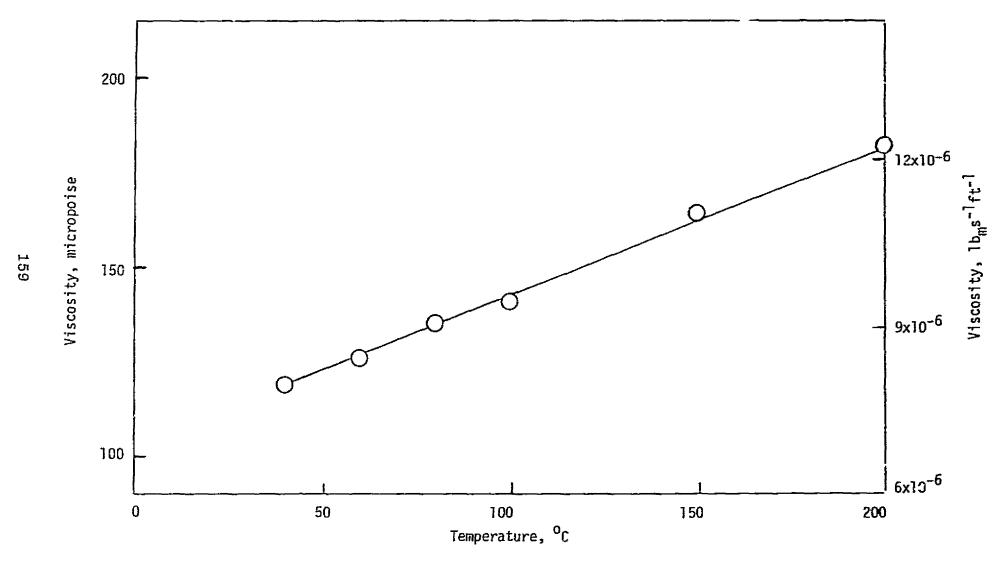


Figure 2.8-4 The Viscosity of Gaseous Dichlorosilane as a Function of Temperature

TABLE 2.8-3

Gaseous Viscosity of Tetrafluorosilane

_amperamme	Viscosity		
°C	micropoise	Nsm ⁻²	lbms-lft-1
40	169.5	16.96 x 10 ⁻⁶	11.40 X 10 ⁻⁶
60	180.7	18.07 x 10 ⁻⁶	12.14 X 10 ⁻⁶
100	191.7	19 17 x 10 ⁻⁶	12.88 X 10 ⁻⁶
150	208.3	20.83 x 10 ⁻⁶	14.00 x 10 ⁻⁶
200	231.2	23.12 x 10 ⁻⁶	15.54 x 10 ⁻⁶

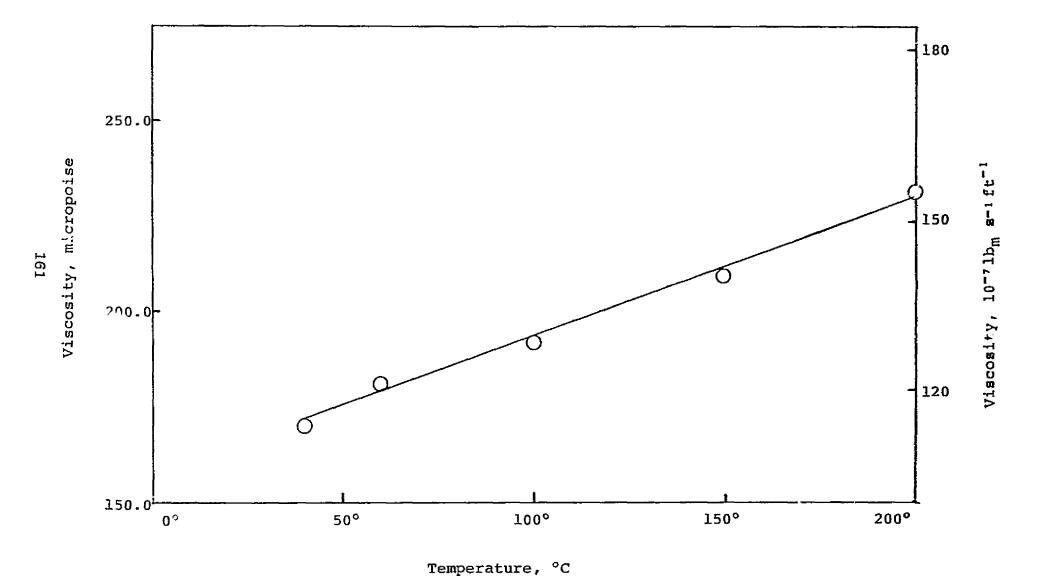


Figure 2.8-5 Viscosity of Gaseous Tetrafluorosilane

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2.9 Silicon Tetrafluoride Generation

Investigations were conducted toward developing a method to genrate silicon tetrafluoride (SiF₄) from an aqueous solution of hexafluorosilicic acid (H₂SiF₆), which is readily available as a product of the phosphate fertilizer industry. The method investigated involved the precipitation of an insoluble salt of hexafluorosilic acid followed by the thermal decomposition of the salt to produce ${\rm SiF_4}$.

Experiments were conducted in which concentrated aqueous solutions of various salts (NaCl, NaF, NaOH, Na₂CO₂) were reacted with a 23% aqueous solution of $\rm H_2SiF_6$ at room temperature:

2 NaX ·
$$H_2SiF_6$$
 · Na_2SiF_6 + 2 HX
where X = C1, F, OH or CO_3

Reaction under these conditions resulted in the immediate formation of a precipitate which could be readily filtered and dried. With each salt, several reactions were carried out with differing stoichiometric amounts of the reactant in order to determine the reactant ratio which would give maximum precipitation of Na₂SiF₆. The results of these investigations are shown in Figures 2.9-1 through 2.9-4.

Figure 2.9-1 shows that the precipitation of Na.SiF₆ with NaCl solutions gives maximum recovery of the SiF₄ recursor at a reactant ratio of slightly greater than 1:1. Increasing the amount of NaCl does not improve the yield any further. The percent yield of Na₂SiF₆ never rises above the 90-95% region due to its slight solubility in water. The precipitate formed was shown to be Na₂SiF₆ by comparing its infrared spectrum with that of an authentic sample. Air drying of the precipitate was shown to leave approximately 1-2% water.

Figure 2.9-2 shows that precipitation of Na₂SiF₆ with NaF solution gives essentially the same results as was obtained with NaCl up to about 1.25:1 reactant ratio. At higher ratios the calculated percent yield of Na₂SiF₆ rises above 100% which indicates that something else is occurring other than the precipitation of Na₂SiF₆. Hydrofluoric acid (HF) is a byproduct of this reaction and will form an insoluble adduct (NaF·HF) which results in the greater than 100% calculated yield.

Precipitation using NaOH solutions (Figure 2.9-3) gives completely different results than that obtained with either NaCl and NaF. The calculated yields (based on $\rm Na_2SiF_6$) are much above 100% and continue to rise up to a reactant ratio of 3:1. The use of NaOH (a strongly basic reagent) resulted in the hydrolysis of the Si-F bond as well as precipitation of $\rm Na_2SiF_6$. The hydrolysis of $\rm SiF_6$ under basic conditions

results in the formation of silicic acids which are hydrated oxides of silicon and these precipitate along with Na₂SiF₆. The formation of silicic acids in these reactions was confirmed by infrared analysis of the precipitate which showed characteristic absorptions for both silicic acid and Na₂SiF₆.

Figure 2.9-4 shows that reaction with Na_2CO_3 (a weakly basic reagent) gives results comparable to those obtained with NaCl solutions. Infrared analysis of the predictate however, indicated the presence of some silicic acids which showed that hydrolysis was occurring with Na_2CO_3 as well as with NaOH.

Studies were also conducted to determine conditions necessary for the efficient precipitation of BaS:F₆ (another possible precursor for the geration of SiF₄). The use of BaS:F₆ as an alternate precursor to SiF₄ (instead of Na₅SiF₆) was investigated because of several advantages this route may have. BaSiF₆ is less soluble than Na₅SiF₆ by a factor of about 30 which should result in a more efficient recovery of SiF₄ from a solution of H₂SiF₆. Secondly, if BaF₂ is used to precipitate the BaSiF₆ there should be no co-precipitation of an adduct with HF (use of NaF in the precipitation of Na₅SiF₆ resulted in the co-precipitation of NaF*HF). Also the Thermal decomposition of BaSiF₆ to give SiF₄ may occur at a lower temperature than that observed for Na₂SiF₆.

The precipitation of BaSiF, was effected by the reaction of a 23% aqueous solution of $\rm H_2SiF_6$ with aqueous solutions of either BaCl $_2$ or BaF $_2$. An immediate precipitate was formed in both

$$BaX_2 + H_2SiF_6 + BaSiF_6 + + 2 HX$$
$$X = C1 \cdot F$$

cases which could be readily tiltered and dried. With each salt, several reactions were carried out with differing stoichiometric amounts of the reactant in order to determine the reactant ratio which would give maximum precipitation of BaSiF_4 . The results of these investigations are presented in Figures 2.9-5 and 2.9-6.

Figure 2.9-5 shows that the precipitation of ${\rm BaStF_6}$ with ${\rm BaCl_9}$ solutions at room temperature gives maximum recovery of the ${\rm SiF_4}$ precursor at a reactant ratio of 1.1. Increasing the amount of ${\rm BaCl_9}$ does not increase the yield any further and the percent yield never rises above about 97% due to the slight solubility of ${\rm BaSiF_6}$.

Figur. 2.9-6 shows the centrs of the reactions of BaF_2 solutions with H_2SiF_6 at $a0^{\circ}C$. Maximum recovery of $BaSiF_6$ is shown to be at a reactant ratio of 1.5 to 1 with the percent yield about 95°. The higher reactant ratio necessary

for efficient precipitation and slightly lower yields than were observed in the reactions with BaCl₂ is due to the low solubility of BaF₂ which dictated the use of considerably larger volumes of water.

Investigations were conducted for the generation of silicon tetrafluoride (SiF₄) by the thermal decomposition of sodium hexafluorosilicate $^4(Na_2SiF_6)$:

$$Na_2SiF_6 \stackrel{\Delta}{\leftarrow} SiF_4 + 2 NaF$$
 (4)

Parameters such as temperature, reaction time, and general reaction conditions were examined in order to determine optimum conditions for efficient SiF_A generation.

Anhydrous sample of Na₂SiF₆ were placed in a quartz tube and heated under various fraction conditions. The amount of SiF_4 generated was determined from the weight loss of the sample after heating based on the stoichiometry in equation 4.

Initially the samples were heated for 1 hour in a closed system of inert gas (N_c) maintained at 1 atmosphere and at constant temperatures ranging from 350°C to 600°C. In no instance was the generation of SiF₄ above 12%. This low yield of the SiF₄ was due to the fact that the thermal decomposition reaction is an equilibrium reaction (eq. 4) and when the decomposition is carried out under conditions which allow an equilibrium to be established, the reaction will proceed no further than the equilibrium point.

As a result of these initial results, the reaction conditions were altered such that a slow stream of N_2 (approximately 200 cc/min) was continuously passed over the sample during the decomposition. Figure 2.9-7 shows the results of these experiments. Heating the samples for 1 hour gave low yields of SiF_4 increased rapidly such that the generation of SiF_4 was essentially quantitative above 550°C.

Based on the above results, it is obvious that the decomposition of Na₂SiF₆ to generate SiF₄ occurs extensively at temporatures above 500°C. Since this data was obtained by heating the samples for an extended period (1 hour at each temperature), the per cent generation of SiF₄ was obtained as a function of reaction time in order to determine the minimum amount of heating required to produce high yields of SiF₄. Figures 2.9-8, 2.9-9, 2,9-10 show the results of this type investigation at 500°C, 550°C, and 600°C. Figure 2.9-8 indicates that decomposition is not complete at 500°C even upon neating for a period of 1 hour. At 550°C the generation of SiF₄ approaches completion in 30 minutes (figure 2.9-9) and at 600°C the reaction is essentially complete in 15 minutes.

The generation of silicon tetrafluoride (SiF_4) by the thermal decomposition of barium hexafluorosilicate ($BaSiF_6$) according to equation 9 was investigated. Parameters such

$$BaSiF_6 \stackrel{\Delta}{\rightleftharpoons} SiF_4^{\dagger} + BaF_2 \tag{9}$$

as temperature, reaction time, and general reaction conditions were examined in order to determine optimum conditions for efficient SiF_4 generation by this method.

Samples of anhydrous BaSiF, were placed in a quartz tube and heated at various temperatures for a period of one hour during which tine a slow stream of N_2 (approximately 200 cc/min) was continuously passed over the sample. The amount of SiF_4 generated was determined from the weight loss of the sample after heating based on the stoichiometry in equation 9. The results of these experiments are shown in figure 2.9-11. Heating for 1 hour at temperatures up to about 400°C gave low yields of SiF_4 . At temperatures above 400°C however, nearly quantitative yields were obtained.

Based on the above results, it can be seen that the decomposition of BaSiF, to generate SiF, occurs extensively at temperatures above 400°C. Since this data was obtained by heating the samples for an extended period (1 hour at each temperature), the per cent yield of SiF, was obtained as a function of time in order to determine the minimum amount of heating required to produce high yields of SiF, Figures 2.9-12, 2.9-13, 2.9-14, 2.9-15 show the results of this type investigation at 400°C, 450°C, 500°C, and 550°C respectively. Figure 2.9-12 shows that at 400°C the decomposition of BaSiF, does not approach completion until about 1 hour heating time. Figure 2.9-13 shows that decomposition of BaSiF, is essentially complete after 30 minutes heating at 450°C and figures 2.9-14 & 2.9-15 show that decomposition is complete after only a few minutes (5-10 minutes) at 500°C and 550°C.

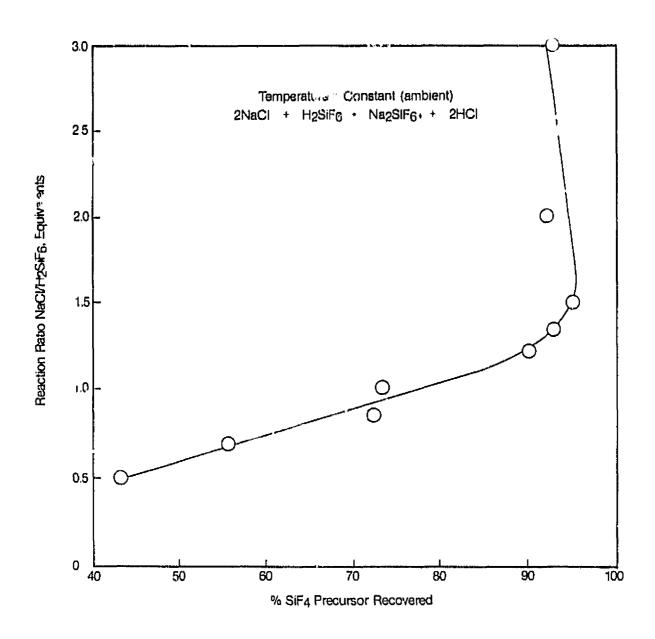


Figure 2.9-1 Variation of SiF4 Precursor Recovery with Reaction Ratio (NaCl Reaction)

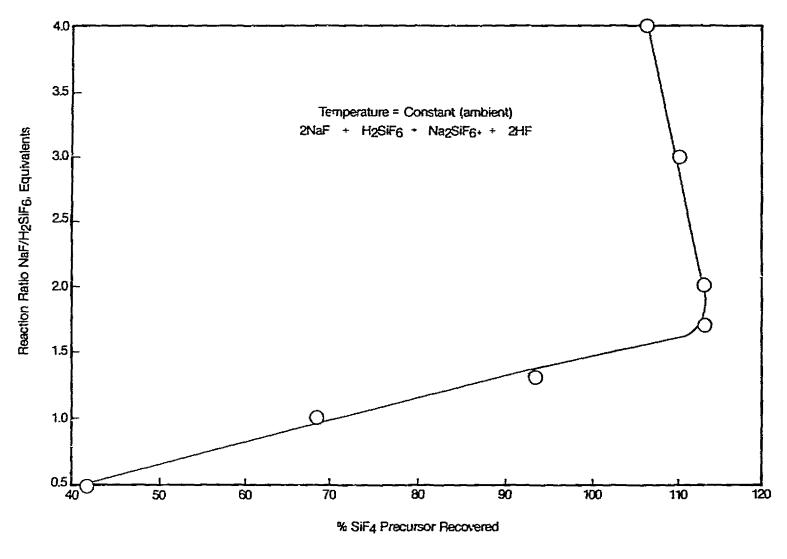


Figure 2.9-2 Variation of SiF4 Precursor Recovery with Reaction Ratio (NaF Reaction)

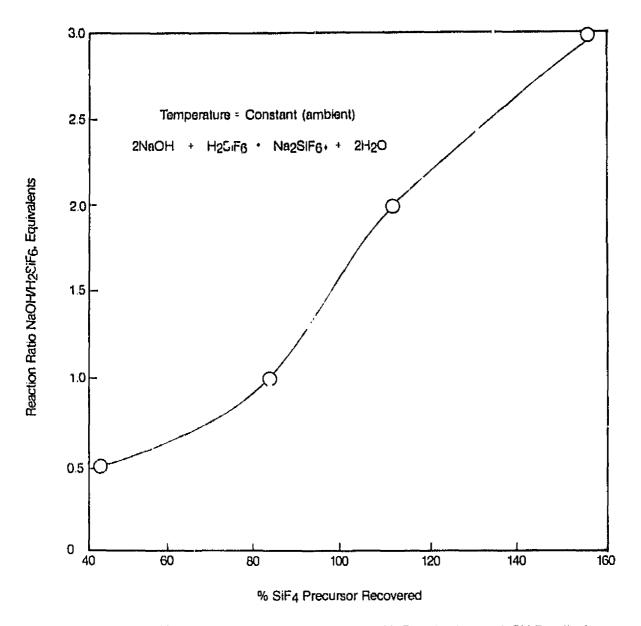


Figure 2.9-3 Variation of SiF4 Precursor Recovery with Reaction Ratio (NaOH Reaction)

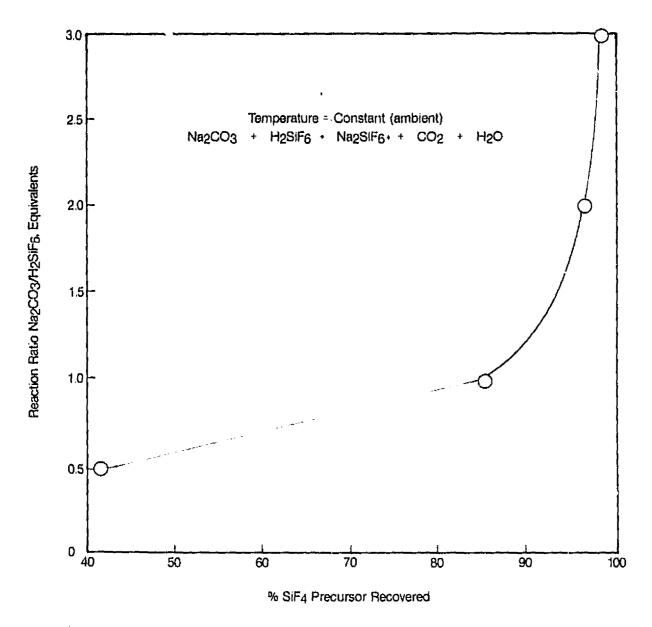


Figure 2.9-4 Variation of SiF4 Precursor Recovery with Reaction Ratio (Na2CO₃ Reaction)

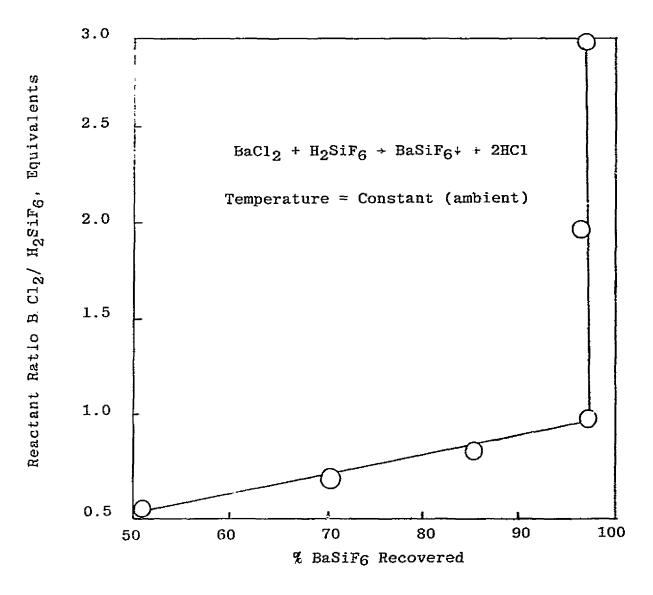


Figure 2.9-5 Variation of BaSiF6 Recovery with Reactant Ratio (BaCl2 Reaction)

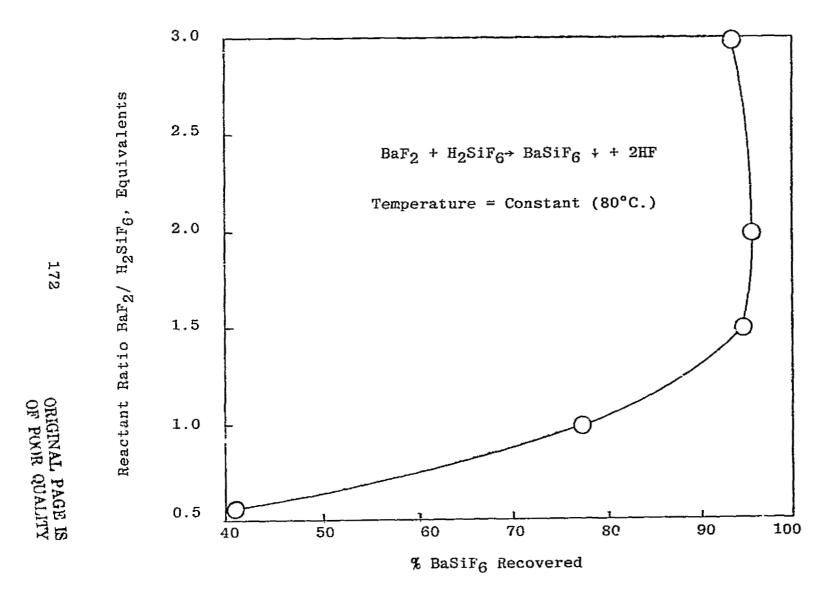


Figure 9 D & Variation of Ragir. December with December Datio (Date to the

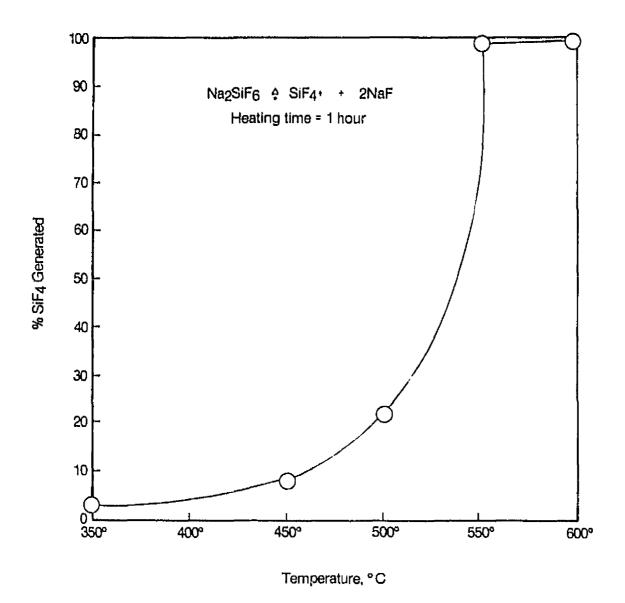


Figure 2.9-7 Variation of % SiF4 Generated with Temperature

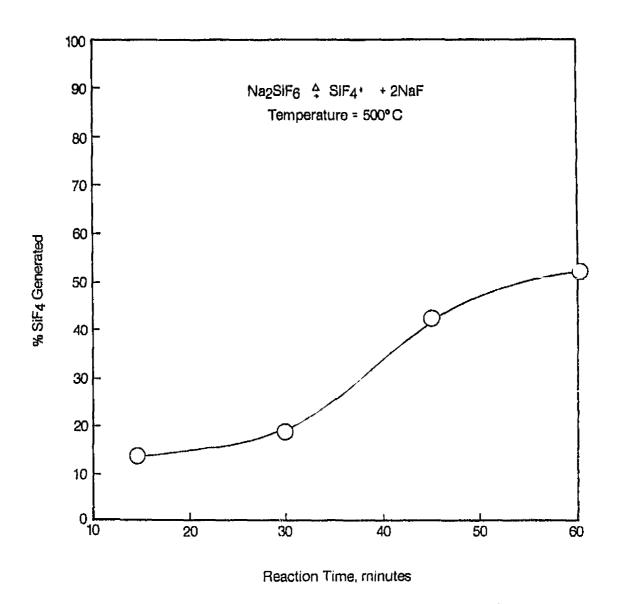


Figure 2.9-8 Variation of % SiF4 Generated with Reaction Time

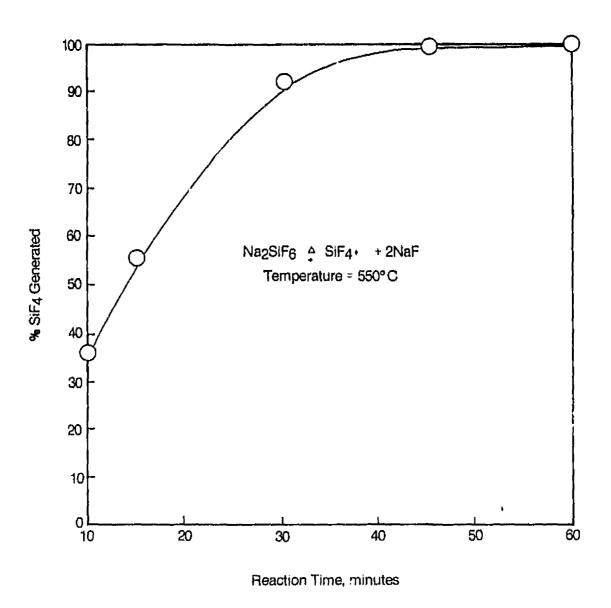


Figure 2.9-9 Variation of % SiF4 Generated with Reaction Time

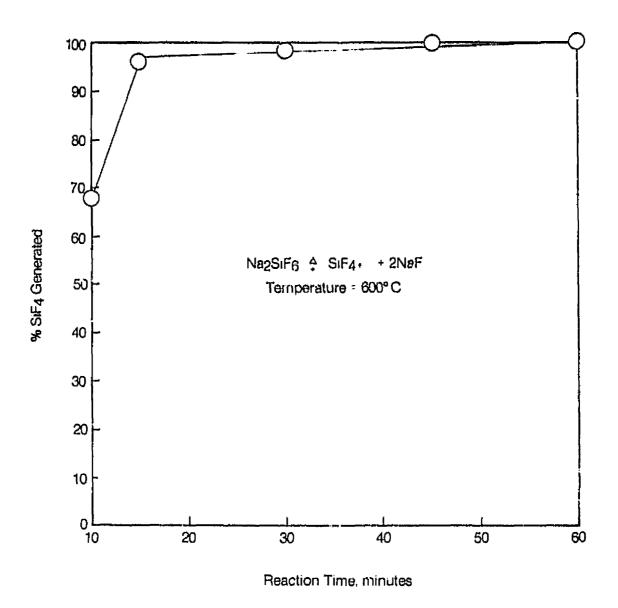


Figure 2.9-10 Variation of % SiF4 Generated with Reaction Time

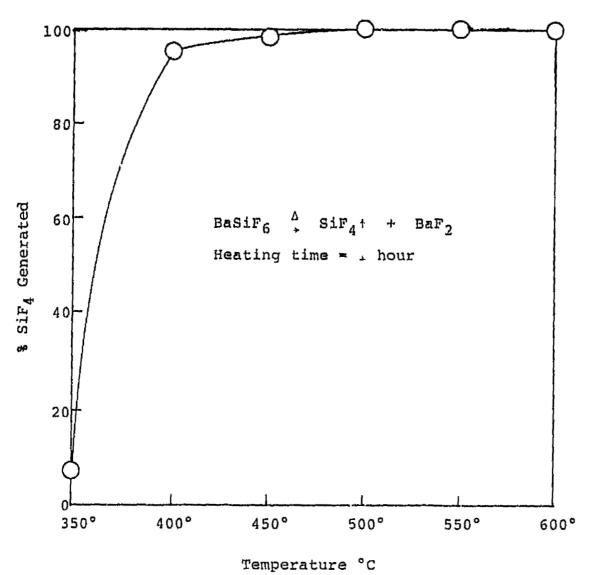


Figure 2.9-11 Variation of % SiF4 Generated with Temperature

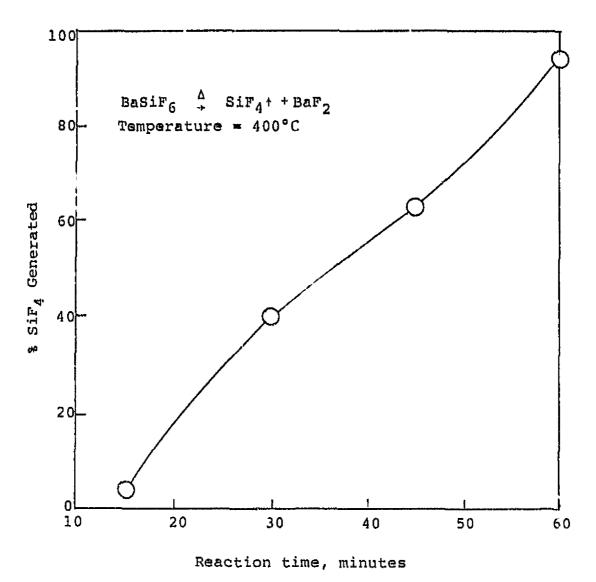


Figure 2.9-12 Variation of % SiF_4 Generated with Reaction Time

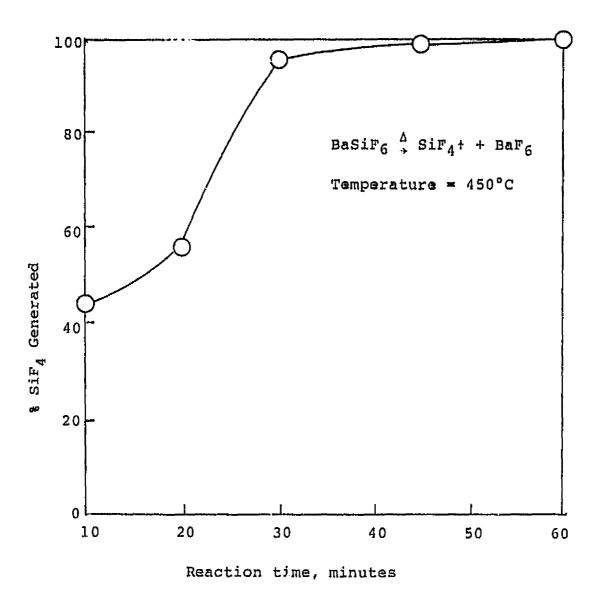


Figure 2.9-13 Variation of % SiF_4 Generated with Reaction Time

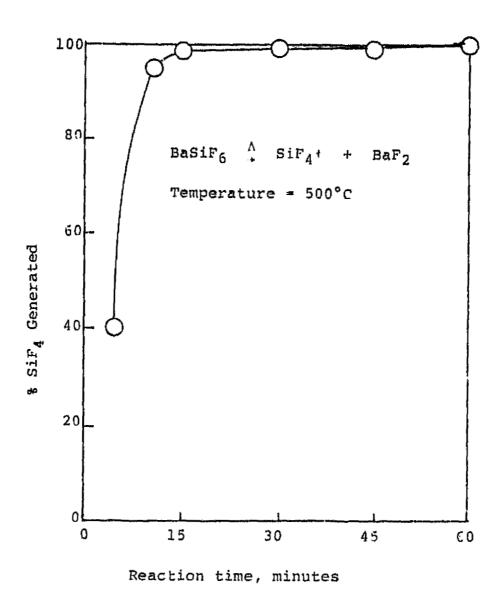


Figure 2.9-14 Variation of % SiF4 Generated with Reaction Time

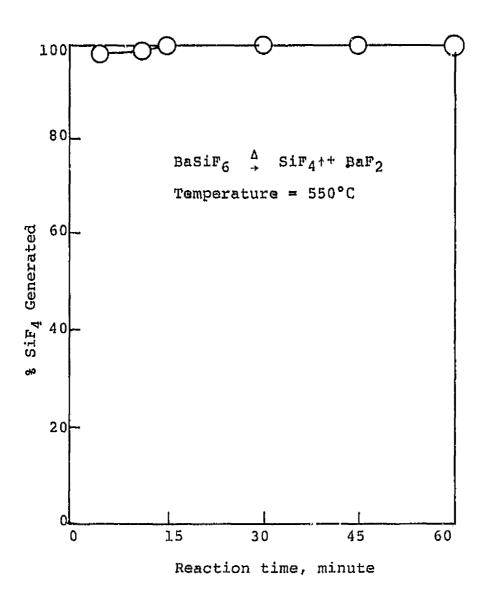


Figure 2.9-15 Variation of % SiF_4 Generated with Reaction Time

3. CHEMICAL ENGINEERING ANALYSES

3.1 Sil Decomposition Process

The chemical engineering analysis activity involves a preliminary process design of a plant to produce silicon via the technology under consideration.

The process flow-diagram for the SiI_4 decomposition process is shown in Figure 3.1-1. This process involves several major processing operations such as fluidization, distillation, condensation, vaporization and deposition unit.

At the beginning of the process, metallurgical grade silicon (M.G. Si) is reacted with iodine (I_2) in a fluidized bed reactor (100°K) to produce silicon tetraiodide. This gas product is condensed and then purified by distillation process.

This purified SiI_4 is vaporized and introduced into a silicon rod reactor where silicon is deposed according to the following reaction

$$SiI_4 \longrightarrow Si + 2I_2$$
 (3.1-1)

The reaction temperature is kept at 1300°K.

Unreacted SiI $_4$ and iodine are condensed and separated by distillation for recycle purpose.

A process design was performed to obtain data for the cost analysis. The design was based on a plant for the production of 1000 metric tons paragraph of polysilicon via this ${\rm SiI}_4$ decomposition process.

The detailed status sheet for the process design package is shown in Table 3.1-1 and is representative of the various sub-items that make up the activity. The summarized results for the preliminary process design are presented in a tabular format to make it easier to locate items of specific interest. The guide for these tables is given below

Base Case Conditions	Ta^{i} Ac	3.1-2
Reaction chemistry	-Table	3 1-3
Raw Material Requirements	-Table	3.1-4
Utility Requirements	-Table	3.1-5
Major Process Equipment	$\operatorname{Teta}_{i}(x) \leftarrow$	3 1-6
Production Labor Requirements-	Table	3.1-7

The process design provides detailed data for raw materials, utilities major or a equipment and production labor requirements which are necessary for polysilicon production

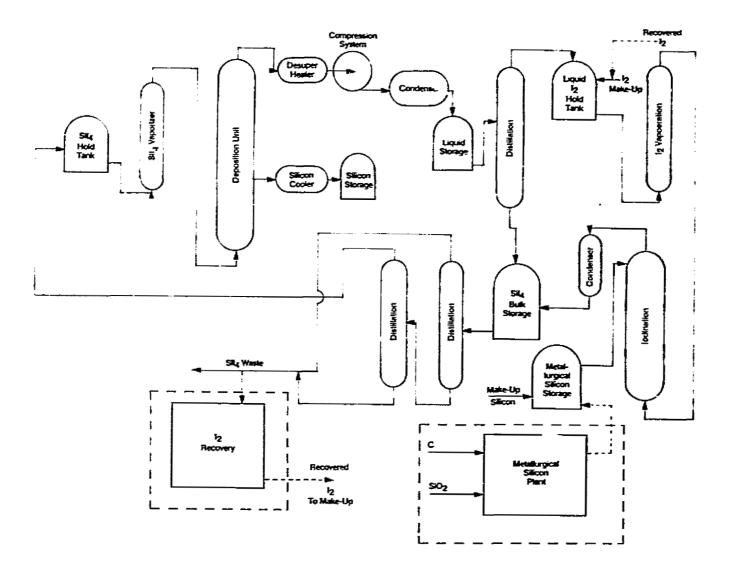


Figure 3.1-1 Process Flow Diagram for SiI₄ Decomposition Process (Battelle)

TABLE 3.1-1 CHEMICAL ENGINEERING ANALYSES: FRELIMINARY PROCESS DESIGN ACTIVITIES FOR SII4 DECOMPOSITION PROCESS

	Prel. Process Design Activity	Status	Prel. Process Design Activity Status
1.	Specify Base Case Conditions 1. Plant Size	•	7. Equipment Design Calculations
	2. Product Specifics		1. Storage Vessels 2. Unit Operations Equipment
		Š	3. Process Data (P, T, rate, etc.)
	3. Additional Conditions	•	4. Additional
2.	Define Reaction Chemistry	•	
	Reactants Products	•	8. List of Major Process Equipment
	2. Equilibrium	•	1. Size
	-		2. Type
3.	Process flow Diagram	•	3. Materials of Construction •
	1. Flow 3-quence, Unit Operations	•	
	2. Process Conditions (T, 2 stc.)	•	8a. Major Technical Factors
	3. Environmental	•	(Potential Problem Areas)
	Company Literaction	•	1. Materials Compatibility
	(Technology Exchange)		2. Process Conditions Limitations
			3. Additional
4.	Material Balance Calculations	•	
	. Raw Materials	•	9. Production Labor Requirements
	Products	•	1. Process Technology
	. By-Product:	•	2. Production Volume
5.	Fnergy Balance Calculations	•	10. Forward for Economic Analysis
_	. Heating	•	
	2. Cooling	•	
	. Additional	•	
6.	Property Data	•	0 Plan
	1. Physical	•	€ In Progress
	2. Thermodynamic	•	● Complete
	3. Additional	•	

TABLE 3.1-2

BASE CASE CONDITIONS FOR SII, DECOMPOSITION PROCESS

- 1. Plant Size
 - Production of 1000 metric tons/year
 - Solar Cell Grade Silicon
- 2. Iodination Reaction
 - Metallurgical grade sili.on and iodine to produce Sild
 - ~ Atmospheric, 1000°K
 - 100% conversion (thermodynamic equilibrium)
 - Fluidized bed
- 3. Sil Purification
 - Recycled and manufactured SiI distilled
 - 10% waste (5% light, 5% heavies)
 - 90% product (heartcut)
- 4. SiI₄ Decomposition
 - Silicon rod reactor, deposition
 - .001 ATM, 1300°K
 - 59,67% conversion (thermodynamic equilibrium)
- 5. SiI_A/I , Recycle
 - Separated by distillation
 - SiI_4 to purification and decomposition
 - I, to iodination
- 6. Operating Ratio
 - Approximately 80% utilization (79.3%)
 - Approximately 7000 hr/year production
- 7. Recovery of Waste SiI₄- Wet tecovery of todine from SiI₄ wastes
 - 90% recovery
 - \$.70 pound of I, recovery costs Recycle I, to I, makeup
- S. Storage Considerations
 - Feed materials (two week supply)
 - Product (two week supply)
 - Process (several days)

1. Silicon Deposition

2. Iodination Reaction

TABLE 3.1-4

RAW MATERIAL REQUIREMENTS FOR SII4 DECOMPOSITION PROCESS

	Raw Material	Requirement lb/Kg of Silicon
1.	Metallurgical Grade Silicon	2.6194
2.	Iodine	7.4954

TABLE 3.1-5

UTILITY REQUIREMENTS FOR ${ m SiI}_4$ DECOMPOSITION PROCESS

Utility/Function	K watt-hours/ Kg of Silicon Product*
Heating and Cooling (10% losses)	27.55
Compressor Train	1.059
Radiant Losses from Deposition	190.
	218.61

^{*} All utility requirements calculated as electricity. Actual useage would involve cooling water, steam, etc.

TABLE 3.1-6 LIST OF MAJOR PROCESS EQUIPMENT FOR SiI_4 DECOMPOSITION PROCESS

		Item	Function	Duty	Size	Material
	I.	Purified SiI 4 Hold Tank	Storage of purified SiI ₄ feed to deposition	One Week	6,254 x 10 gallons	316 s.s.
	2.	Liquid Storage	Cooled overheads from deposition	One Week	5.6 x 10 ⁴ gallons	316 s.s.
	3.	Silicon Product Storage	Product for sales	Two Weeks	1.097×10^4 gallons	C.s.
18	4.	Liquid I ₂ Storage	I ₂ separated from SiI ₄	One Week	2.962 x 10 ⁴ gallons	316 s.s.
189-199	5,	SiI ₄ Bulk Storage	SiI ₄ separated from I ₂	One Week	6.948 x 10 ⁴ gallons	316 s.s.
	6.	Metallurgical Silicon Storage	Raw material storage for manufacturing	Two Weeks	1.304 x 10 ⁴ gallons	c.s.
	7.	Feed Tank	Purification column 2 feed tank	Eight Hours	4.008 x 10 ³ gallons	316 S.S.
	8.	SiI ₄ Vaporizer	Vaporize SiI ₄ for deposition unit	+29.86 K cal/gmoles Si produced	121.5 ft ²	C.S. with Hastelloy tubes
	9 .	Silicon Cooler	Cocl product silicon for storage and ship- ment	-5.7 K cal/gmole Si	1D = 1.85 ft L = 5.67 ft	6 units - Graphii or Quartz
	10.	Deposition Condenser	Condense overheads for recycle	-37.146 K cal/gmoles Si	1510.8 ft ²	C.S. with Hastelloy tubes

TABLE 3.1-6 (Continued)

	Item	Function	Duty	Size	Material
11.	Separation Column Preheater	Preheat feed to bubble point for dist.	+3.289 K cal/gmmole Si	22.3 ft ²	C.S. with Hastelloy tribes
12.	Separation Column O/H Condenser	Provide liquid reflux to column	-36.66	216.2 ft ²	C.S. básh Hastelloy tubes
13.	Separation Column Calandria	Provide vapor rate to column	÷42.86	210 ft ²	C.S. with Hastelloy tubes
14.	Separation Column O/H After Cooler	Cool I ₂ at B.P. to 130°C for storage	- 2.1	14.24 ft ²	C.S. with Hastelloy tubes
15.	I ₂ Vaporizer	Vaporize I ₂ for iodination unit	+26.71	113 ft ²	C.S. with Hastelloy tubes
16.	Iodination O/H Condenser	Condense bulk SiI ₄ vapors from iodina- tion	-33.aS	1377.42 ft ²	C.S. with Hastelloy tubes
17.	Separation Column Bottoms After Cooler	Cool SiI ₄ at B.P. to 150°C for storage	~ 4. 2	28.48 ft ²	C.S. with Hastelloy tubes
18.	Tet Purification Preheater	Bring SiI ₄ to bubble point for distillation	+11.41	77.38 ft ²	C.S. with Hastelloy tribes
19.	Purification Column 1 O/H Condenser	Provide reflux for operation of column	-54.42	221 ft ²	C.S. with Hastelloy tubes
20.	Purification Column l Calandria	Provide vapor for column operation	54.42	221 ft ²	C.S. with Hastellow tubes
21.	Purification Column 2 O/H Condenser	Provide reflux for operation of column	-52.24	212 ft ²	C.S. with Hastelloy tebes

		Item	<u>Punction</u>	Duty	Size	Material
	22.	Purification Column 2 Calandria	Provide vapor for operation of column	52.24 K cal/gmole Si	212 ft ²	C.S. with Eastelloy tubes
	23.	Purification After Cooler	Cool purified SiI ₄ to 150°C for storage	-10,27	69.7 ft ²	C.S. with Hastelloy tubes
	24.	De-superheater	Cool for compression	-24.03	163 ft ² each	6 units - Quartz or Graphite
	25.	Purified SiI ₄ Pump	Feed to SiI ₄ vaporizer	6.2 дра	100 ft of head	316 s.s.
	26.	Deposition Com- pressor(s)	Return deposition gases to atmospheric pressure	-23.03 K cal/gmole 4.155 x 10 ⁵ ft ³ /min	184.1 horsepower	316 S.S.
	27.	I ₂ /SiI ₄ Liquid Pump	Pump to I ₂ /SiI ₄ separation	5.56 gp a	100 fc of head	316 S.S.
201	28.	I ₂ /SiI ₄ Separa- tion Column Over- heads Pump	Pump O/H I ₂ for reflux and storage	5.66 gpm	100 ft	316 s.s.
	29.	I ₂ /SiI ₄ Separa- tion Column Bottoms Pump	Pump bottoms for reboil and SiI_4 storage	18.‡िक्सच	150 ft	316 s.s.
	30.	I ₂ Pump	Pump liquid I ₂ through vaporization and iodination	3.54 gg#	150 ft	316 s.s.
	31.	SiI _A Pump	Pump liquid SiI ₄ to purification	6.89 qom	100 ft	316 S.S.
	32.	Tet Purification Column 1 O/H Pump	Pump O/H to reflux and waste	19.75 gpm	190 ft	316 S.S.

TABLE 3.1-6 (Continued)

		Item	Function	Duty	Size	Meterial
	33.	Tet Purification Column 1 Bottoms Pump	Pump bottoms for reboil and remove product	27.25 дря	150 ft	316 s.s.
	34.	Tet Purification Column 2 Feed Pump	Feed to Column 2	°.504 ggm.	50 ft	316 S.S.
	35.	Tet Purification Column 2 O/H Pump	Pump for reflux and to purified stage	I.E. SHIT GERMA	100 ft	316 s.s.
	36.	Tet Purification Column 2 Bottoms Pummp	Pump bottoms for repoil and waste	19.35 çom	150 ft	316 s.s.
	37.	Deposition Unit	Produce Si from SiI ₄ + Si + 2I ₂	144 Kg of Si/hour + 89.82 K cal/gmole Si	6 at 1018 ft ² each	Quartz
202	38.	Iodination Reactor	Produce SiI ₄ from met grade Si. Si + 2I ₂ → SiI ₄	3263.25 Kg/hr of raw SiI ₄ , -44.93 K cal/gmoles	23 inch ID by 10 ft tall	Graphite
	39.	I ₂ /SiI ₄ Distilla- tion Column	Separate I ₂ from SiI ₄ for recycle	Separate 1903.36 Kg/hr SiI ₄ + 2602.61 Kg/hr I ₂	20 ft by 23 inch ID	316 s.s.
	40.	Purification Column 1	Purify SiI ₄ by 5% cut off top to waste	Feed rate 5166.61 Kg/hr SiI ₄	47.6 ft by 26.5 inch ID	316 s.s.
	41.	Furification Column 2	Purify SiI _A by 5% cut off bottom to waste	Feed rate 4908.28 Kg/nr SiI ₄	23.8 ft by 26 inch IU	316 s.s.

TABLE 3.1-7 PRODUCTION LABOR REQUIREMENTS
OF SiI₄ DECOMPOSITION PROCESS

	Unit Operation	Type	Skilled Man Hrs/Day Unit	Semiskilled Man Hrs/Day Unit
1.	Vaporization	В	23	
2.	Deposition Unit	A	36	
3.	Compression System	B	23	
4.	Vapor Condensation	В	23	
5.	I2/SiI4 Distillation	* _y ,	16	
6.	Iodinaction	B	23	
7.	Tet Purification	С	16	
8,	Materials Handling	A		36
9.	Product Handling	A	### **********************************	<u>36</u>
			160	72

Skilled: .0584 Man-hrs/KgSi

Semiskilled: .0263 Man-hrs/KgSi
TOTAL .0847 Man-hrs/KgSi

NOTES

- 1. A Batch Process or Multiple Small Units
 - B Average Process
 - C Automated Process
- 2. Manhours/Day Unit from Figure 4-6, Peters & Timmerhaus (7).

3.2 Conventional Process for Polysilicon (Sigmens Technology)

The chemical engineering analysis activity involves a preliminary process design of a plant to produce polysilicon through the conventional process (Siemens Technology).

The process flowsheet for the conventional polysilicon process, consisting of several major processing operations of hydrochlorination, condensation, distillation and chemical vapor deposition, is shown in Figure 3.2-1.

Initially, metallurgical grade silicon (MGSi) is reacted with anhydrous hydrogen chloride (NC1) in a fluidized bed (550-650°K) to produce a mixture of chlorosilanes, which is primarily trichlorosilane (TCS) and silicon tetrachloride (TET). Since the reactions are highly exothermic, heat transfer for removal of heat of reaction is required to maintain reaction temperature control. The mixture of chlorosilanes from the reaction is condensed and subjected to a several stage distillation to separate by-products and remove impurities.

The purified TCS is reacted with hydrogen (H₂) in a rod reactor to obtain polysilicon deposition via the representative reaction:

$$SiHCl_3 + H_2 + Si + 3HCl$$
 (3.2-1)

The deposition reaction occurs on the surface of a hot rod (1000-1100°C) which is heated by passage of electrical current through the rod. Large electrical energy requirements are necessary because of the endothermic reaction, radiation heat losses and incomplete conversion of the TCS. Unreacted chlorosilanes and hydrogen are separated and recycled. Silicon tetrachloride is not recycled.

A process design was performed to obtain data for the cost analysis. The design was based on a plant for the production of 1000 metric tons per year of semiconductor grade polysilicon via the conventional Siemens process.

The detailed status sheet for the process design package is shown in Table 3.2-1 and is representative of the various sub-items that make up the activity. The summarized results for the preliminary process design are presented in a tabular format to make it easier to locate items of specific interest. The guide for these tables is given below:

- · Production Labor Requirements----Table 3.2-7

The process design provides detailed data for raw materials, utilities, major process equpment and production labor requirements which are necessary for polysilicon production.

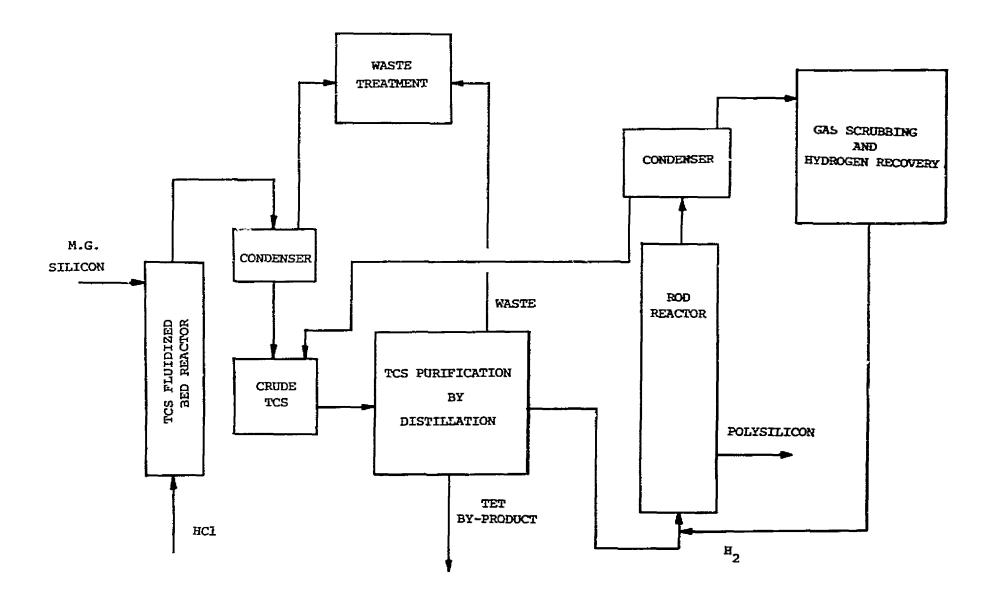


Figure 3.2-1 Preliminary Process Flowsheet for Conventional Polysilicon Process

TABLE 3.2-1 CHEMICAL ENGINEERING ANALYSES: PRELIMINARY PROCESS DESIGN ACTIVITIES FOR CONVENTIONAL POLYSILICON PROCESS

	Prel. Process Design Activity	Status	Prel. Process Design Activity Status
1.	Specify Base Case Conditions 1. Plant Size 2. Product Specifics 3. Additional Conditions		7. Equipment Design Calculations 1. Storage Vessels 2. Unit Operations Equipment 3. Process Data (P, T, rate, etc.) 4. Additional
2.	Define Reaction Chemistry 1. Reactants, Products 2. Equilibrium	•	8. List of Major Process Equipment 1. Size 2. Type
3.	Process Flow Diagram 1. Flow Sequence, Unit Operations 2. Process Conditions (T, P, etc.) 3. Environmental 4. Company Interaction (Technology Exchange)	•	3. Materials of Construction 8a. Major Technical Factors (Potential Problem Areas) 1. Materials Compatibility 2. Process Conditions Limitations 3. Additional
4.	Material Balance Calculations 1. Raw Materials 2. Products 3. By-Products	• • •	9. Production Labor Requirements 1. Process Technology 2. Production Volume
5.	Energy Balance Calculations 1. Heating 2. Cooling 3. Additional	•	10. Forward for Economic Analysis
6.	Property Data 1. Physical 2. Thermodynamic 3. Additional	•	0 Plan 9 In Progress • Complete

TABLE 3.2-2 BASE CASE CONDITIONS FOR CONVENTIONAL POLYSILICON PROCESS

- Plant Size - 1000 metric tons per year - Semiconductor grade silicon 2. Production of TCS - Fluidized Bed, 600 K, low pressure (65 PSIA) - Metallurgical grade silicon plus HCl gas - Chlorosilane content in condensed reator gas by moles (ref. 32) 91.5% TCS (8iCl₃H) 5.2% TET (SiCl₄) 1.4% DCS (SiCl H2) 1.9% Heavies - Slight excess HCl in reator gas (1%) - Hydrogen burned 3. TCS Purification (ref. 31) - Distillation - 5% lights to waste (5% of TCS & TET) - Separate TCS and TET - 5% heavies from TCS & TET to waste - TET for by-product sales - TCS to rod reactor 4. Silicon Production - Rod reactor at 1050°C, 20 PSIA - Hydrogen to reduce TCS - Entering gas analysis 10% TCS 90% H₂ - 8.17 moles TCS in/mole of S; production in an operating reactor
 - 8.17 moles TCS in/mole of S; production in an operating reactor Exit gas analysis (ref. 20)

4.339% TET 4.457% TCS .089% DCS 2.197% HC1 88.92% H₂

- 5. Waste Treatment
 - Light and heavy cuts from distillation to waste treatment
 - Vapors from TCS reactor condenser to scrubber
 - Vapor from rod reactor to scrubber
 - All waste streams neutralized with NaOH

TABLE 3.2-2 (Continued)

- 6. Recycles
 - H2 from rod reactor dried and returned, 5% losses
 - Chlorosilanes from rod reactor condensed off gas recycled to purification (distillation)
- 7. Operating Ratio
 - Approximately 90% utilization
 - Approximately 7880 hour/year production
- 8. Storage Considerations
 - Feed materials (two week supply)
 - Product (two week supply)
 - Process (several days)
- 9. Filament Pullers
 - Pull rate of 50-100 inches/hour
 - Average of 72 inches/hour used
 - 1/4" Filaments for silicon deposition needed

TABLE 3.2-3

REACTION CHEMISTRY FOR CONVENTIONAL POLYSILICON PROCESS

1. TCS Reactor

$$si + 3 HC1 \rightarrow siHC1_3 + H_2$$

 $si + 4 HC1 \rightarrow siC1_4 + 2H_2$
 $si + 2HC1 \rightarrow siH_2C1_2$

2. Rod Reactor

$$sihcl_3 + H_2 + si + 3HC1$$

 $sihcl_3 + HC1 + sicl_4 + H_2$
 $sihcl_3 + H_2 + siH_2Cl_2 + HC1$

3. Waste Treatment

$$\begin{array}{l} {\rm sihCl}_3 + 2{\rm H}_2{\rm O} + {\rm sio}_2 + 3{\rm HCl} + {\rm H}_2 \\ {\rm sicl}_4 + 2{\rm H}_2{\rm O} + {\rm sio}_2 + 4{\rm HCl} \\ {\rm sih}_2{\rm Cl}_2 + 2{\rm H}_2{\rm O} + {\rm sio}_2 + 2{\rm HCl} + 2{\rm H}_2 \\ {\rm HCl} + {\rm NaOH} + {\rm NaCl} + {\rm H}_2{\rm O} \end{array}$$

TABLE 3.2-4 RAW MATERIAL REQUIREMENTS FOR CONVENTIONAL POLYSILICON PROCESS

	Raw Material	Requirement lb/Kg of Silicon
1.	M. G. Silicon	6.72 Kg/Kg
2.	Anhydrous HCl	57.96
3.	Hydrogen	.828
4.	Caustic (50% NaOH)	53.29
5.	SiCl _A (By Product)	46.12

TABLE 3.2-5

UTILITY REQUIREMENTS FOR CONVENTIONAL POLYSILICON PROCESS

	UTILITY/FUNCTION		REQUIREMENTS/Kg OF SILICON PRODUCT
3 10	Restricity		384.62 Kw-Hr
1 2 3	All pump motors (16 motors) 2. 2 compressor motors 3. Polysilicon Rod Reactor 4. Filament Pullers	(.339) (9.243) (375) (.0244)	
2. s 1 2 3 4 8 6 7 8 10	Rteam (250 PSIA) HCl Vaporiser Caustic Storage Tank H1 Scrubber Vapor Heater H2 Distillation Column Calandria H3 Distillation Column Calandria H3 Distillation Column Calandria TCS Vaporizer H2 Scrubber Vapor Heater Liquid Recycle Heater H4 Distillation Column Calandria Rod Reactor	(7.07) (1.82) (.276) (38.75) (47.73) (25.24) (10.79) (3.4) (5.52)	152 Pounds
1 2 3 4	Cooling Water 1. TCS Reactor Off Gas Cooler 2. Rod Reactor Off Gas Cooler 3. #4 Distillation Column Condenser 4. Polysilicon Rod Reactor Cooling End Plates 5. TCS Reactor Off Gas Compressor 6. Rod Reactor Off Gas Compressor	(13.91) (334) (37.24) (473) (11.12) (115.2)	984.5 Gallons
1	Process Water 1. #2 Gas Scrubber 2. #1 Gas Scrubber 3. To Make Steam In Cooling Rod Reactor Side Walls	(31.36) (134.82) (154.7)	320.9 Gallons
1	Refrigerant (-40 ⁰ F) . TCS Reactor Off Gas Condenser . Rod Reactor Off Gas Condenser	(12.57) (29.52)	42.1 M BTU
) 2	Refrigerant (34°F) 1. #1 Distillation Column Condenser 2. #2 Distillation Column Condenser 3. #3 Distillation Column Condenser	(34) (37.4) (20.85)	92.3 M BTU
1	ligh Temperature Heat Exchange Fluid TCS Fluidized Bed Reactor Nitrogen Heater	(581) (0.61)	582 Pounds
1	Nitrogen L. Molecular Sieves L. Polysilicon Rod Reactor Purge 21	(328.5) (20.64) 12	349.1 SCF

TABLE 3.2-6

LIST OF MAJOR PROCESS EQUIPMENT FOR CONVENTIONAL POLYSILICON PROCESS

Materials

							materials
			Type	Function	Duty	Size	of Construction
	1.	(T1)	M.G. Silicon Storage Hopper	Raw Material Storage	2 Weeks Storage	6.5 x 10 ⁴ gallons	cs
	2.	(T2)	Liquid HCl Storage Tank	Raw Material Storage	2 Weeks Storage	2.5 x 10 ⁵ gallons 250 PSIA	Nickel Steel
	3.	(T3)	Crude TCS Hold Tanks (3)	Peed for Purification	1 Week Storage	2.77 x 10 ⁵ gallons (each)	cs
	4.	(T4)	Waste Hold Tank	Feed Por Waste Treatment	1 Week Storage	3.025 x 10 ⁴ gallons	cs
	5.	(T5)	TCS Reactor Off Gas Flash Tank	Phase Separation		1 ft. in diameter by 4 ft. tall, 300 PSIA	SS
213	6.	(T6)	Hydrogen Storage Tank	Make-up For Losses	8 Hours Backup for Pipeline Failure	7.24 x 10 ⁴ gallons Spherical 250 PSIA	cs
	7.	(T7)	Polysilicon Storage Space	Final Product Storage	2 Weeks Storage	1300 ft. 3 of space	CS
	8.	(T8)	TET Storage Tanks (2)	Final By-product Storage	2 Weeks Storage	1.62 x 10 ⁵ Gallous (each)	cs
	9.	(T9)	TET Feed Tanks (2)	Feed for Distillation Column #4	1 Week S _rage	8.83 x 10 ⁴ Gallons (each)	cs
	10.	(T10)	TCS Feed Tanks (3)	Peed for Distillation Column #3	I Day Storage	2.47 x 10 ⁴ Gallons (each)	cs
	11.	(T11)	TCS Storage Tanks (3)	Purified TCS Hold-Up Feed to Rod Reactor	1 Week Storage	1.64 x 10 ⁵ Gallons (each)	cs
	12.	(T12)	TET/TCS Feed Tanks (3)	Feed for Distillation Column #2	1 Day Storage	3.75 x 10 ⁴ Gallons (each)	cs

	13.	(T13)	Caustic Storage Tank	Raw Material Storage	2 Week Storage 1.91 x 10 ⁵ BTU/HR	1.92 x 10 ⁵ Gallons	ss
	14.	(T14)	#1 Distillation Condenser Flash Tank	Phase Separation		l Pt. in Diameter by 4 Peet Tall	cs
	15.	(T15)	Rod Reactor Off Gas Flash Tank	Phase Separation		1 Pt. in Diameter by 4 Feet Tall 300 PSIA	SS
	16.	(H1)	HC1 Vaporizer	Vaporize Feed To TCS Reactor	7.5 x 10 ⁵ BTU/Er	38.29 Pt. 2 250 PSIA Shell	\$\$/\$ \$
ĸ	17.	(H2)	TCS Reactor Off Gas Cooler	Cool Reaction Gas	4.4 x 10 ⁵ BTU/Hr	224 Ft. 2 65 PSIA Tubes	CS/SS
	18.	(H3)	TCS Reactor Off Gas Condenser	Condense Reaction Gas	1.6 x 10 ⁶ BTU/Hr	1423 Pt. 2 300 PSIA Tubes	SS/SS
	19.	(H 4)	#1 Scrubber Vapor Heater	Heat Vapor Wastes to 40°F for Scrubbing	3 x 10 BTU/Hr	15.7 Ft. ² 250 PSIA Shell	CS/SS
	20.	(H5)	#1 Distillation Column Condenser	Condense Overheads for Relux	4.31 x 10 ⁶ BTU/Hr	1540 Ft. ²	CS/SS
	21.	(H6)	#l Distillation Column Calandria	Reboiler for Column #1	4 x 10 ⁶ BTU/Hr	311. Ft. ² 250 PSIA Shell	CS/SS
	22.	(H7)	#2 Distillation Column Condenser	Condense Overheads For Reflux	4.7 x 10 ⁶ BTU/Hr	1555 Ft. ²	cs/cs
	23.	(8H)	#2 Distillation Column Calandria	Reboiler for Column #2	5 x 10 ⁶ BTU/Hr	402.4 Ft. 250 PSIA Shell	CS/SS
	24.	(H9)	#3 Distillation Column Condenser	Condense Overheads for Reflux	2.64 x 10 ⁶ BTU/Hr	867 Ft. ²	cs/cs

25.	(H10)	#3 Distillation Column Calandria	Reboiler for Column #3	2.64 x 10 ⁶ BTU/Hr	173 Ft. 250 PSIA Shell	CS/SS
26.	(H11)	TCS Vaporizer	Vaporize Feed To Rod Reactor	1.13 x 10 ⁶ BTU/Hr	3 Ft. ² 250 PSTA Shell	cs/cs
27.	(Hl2)	Rod Reactor Off Gas Cooler	Cool Reaction Gas	1.06 x 10 ⁷ BTU/Er	2519 Pt. 2 20 PSIA	CS/SS
28.	(H13)	Rod Reactor Off Gas Condenser	Condense Reaction Gas	3.74 x 10 ⁶ BTU/Hr	3341 Ft. 2 300 PSIA Tubes	SS/ SS
29.	(H14)	#2 Scrubber Vapor Heater	Heat Vapor Wastes to 40°F for Scrubbing	3.56 x 10 ⁵ BTU/Er	180 Ft. ² 250 PSIA Shell	CS/SS
	(H15)	Liquid Recycle Heater	Heat Cold Recycle Liquid (Crude TCS) to 80 ⁰ F for Storage	5.79 x 10 ⁵ BTU/Hr	30.6 Ft. ² 250 PSIA Shell	SS/SS
10 10 13 11.	(H16)	#4 Distillation Column Condenser	Conderser Overheads for Reflux	1.18 x 10 ⁶ BTU/Hr	513 Ft. ²	CS/CS
32.	(H17)	#4 Distillation Column Calandria	Reboiler for Column #4	1.18 x 10 ⁶ BTU/Hr	95 Ft. ² 250 PSIA Shell	CS/SS
33.	(H18)	Nitrogen Heater	Heat Regenerator Gas for Molecular Sieves	2.46 x 10 ⁴ BTU/Hr	44.8 Pt. ²	cs/cs
34.	(PL)	TCS Reactor Off Gas Compressor	Compress Reaction Gas For Condensation	3.52 x 10 ⁵ BTU/Hr	138.2 Horsepower	cs
35.	(P2)	Caustic Supply Pump	Supply Caustic for Waste Neutralization and Gas Scrubbers		9 open 100 Ft. of Head	SS
36.	(P3)	#1 Distillation Column Overheads Pump	Supply Reflux and Remove Waste to Waste Hold Tank		62.2 ggma 100 ?t. of Head	CS*

3.	7. (P4)	#1 Distillation Column Calandria Pump	Forced Convection Pump		93 ogra 150 Ft. of Head	CS*
38	3. (P5)	TET/TCS Feed Pump	Peed #2 Distillation Column		26.1 gpm 100 Pt. of Head	CS*
35). (P6)	#2 Distillation Column Overheads Pump	Supply Relux, Pump Overhead to TCS Feed Tank		70 gpm 100 Ft. of Head	CS*
40). (P7)	TCS Feed Pump	Peed #3 Distillation Column		21 gpm 190 Ft. of Head	C8*
4]	. (P8)	#2 Distillation Column Calandria Pump	Porced Convection Pump		104 ggmm 150 Pt. of Head	C2*
2 42 6	. (P9)	#3 Distillation Column Overhead Pump	Supply Reflux, Pump Overheads to TCS Storage Tank		39 ggmm 100 Pt. of Ps:d	CS *
43	(P10)	Rod Reactor TCS Feed Pump	Feed TCS to Rod Reactor		15 gpm 190 Ft. of Head	CS*
44	. (P11)	#3 Distillation Column Calandria Pump	Porced Convection Pump		39 ggmm 150 Pt. of Head	C5*
45	. (P12)	Rod Reactor Off Gas Compressor	Compress Reaction Gas for Condensation	3.65 x 10 ⁶ BTU/Hr	1434 Horsepower	cs
46	. (PI3)	#4 Distillation Column Overheads Pump	Supply Reflux Pump TET by product to TET Storage Tank		21.59 ggmm 100 Pt. of Head	CS*
47	. (P14)	#4 Distillation Column Calandria Pump	Forced Convection Pump		22.4 gpm 100 Ft. of Bead	CS*

NOTES

^{*}Includes incremental higher cost for special purity requirements.

48.	(P15)	TET Feed Pump	Peed #4 Distillation Column		9.2 gom 100 Pt. of Head	CS*
49.	(P16)	Waste Treatment Pump	Pump from Waste Hold To Waste Treatment		2.8 gpm 50 Pt. of Head	CS
50.	(PI7)	Crude TCS Feed Pump	Feed Purification Area		28 ggm 100 Pt. of Head	CS*
51.	(PI8)	Process Water Feed Pump	Feed Process Water to Scrubber and Waste Trestment		350 gpm 190 Ft. of Head	cs
52.	(C1)	*1 Gas Scrubber	Scrub Gas Wastes from TCS Practor Off Gas		43 Ft. Tall D = 3½ Ft.	SS
53.	(C2)	#2 Gas Scrubber	Scrub Gas Wastes from H16, H3, H5		40 Ft. Tall D = 2½ Ft.	SS
1254. 7	(C3)	#1 Distillation Column	Separate Light Impurities to Maste		29 Trays 24 inches apart 3 3/4 Ft. in Diameter	cs
55.	(C4)	#2 Distillation Column	Separate TET and TCS		29 Trays 24 inches apart 4½ Pt. in Diameter	cs
56.	(CS)	#3 Discillation Column	Separate Heavies TCS to Waste		15 Trays 20 inches apart 3 Ft. in diameter	cs
57.	(C6)	#4 Distillation Column	Separate Heavies TET to Waste		15 Trays 20 inches apart 2½ Feet in Diametez	CS
58.	(Rl)	TCS Fluidized Bed Reactor	Production of TCS For Rod Reactor	4.552 x 10 ⁶ BTU/hr (Cooling)	D = 2.61 it. L = 28.8 Pt. 64, 1° O D Cooling Tubes 9.4° Long	SS

59. (R2)	Polysilicon Rod Reactors (305)	Production of Polysilicon		Hairpin Reactor (2 hair- pins, 3 Ft. long, 6 Inch Dia.)	Quartz
60. (Al)	Molecular Sieves (2)	Dry Out Rod Reactor Off Gas For Hydrogen Recycle		D = 3.5 Pt. L = 14.4 Pt.	cs
61. (A2)	Fines Separator	Remove Solids From Flu. lized Bed Reactor Off Gas		12" Cyclone Separator	SS
62. (A3)	Hydrogen Flare	Dispose of Hydrogen Produced in TCS Fluidized Beá Reactor	8.94 x 10 ⁶ BTU/Hr	30 Peet High Stack 6" diameter	cs
63. (A4) 22 18	Filament Pullers	Production of 1/4" filament Polysilicon depositon	s for		

TABLE 3.2-7

PRODUCTION LABOR REQUIREMENTS FOR CONVENTIONAL POLYSILICON PROCESS

			Skilled La	bor	Semiskil:	led Labor
	Unit Operation	Type	Man Hrs/Day F	er Kg Si	Per Day	Per Kg Si
1.	PCS Production	A	80	.0292		
2.	Vaporization	В	60	.0219		
3.	Vapor Compression	В	61	.0219		
4.	Vapor Condensation	В	60	.0219		
5.	TCS/TET Separation	C	40	.0146		
6.	TCS Purification	С	35	.0128		
7.	TET Purification	C	30	.011		
8.	Filament Pullers		120	.0438		
9.	Gas Scrubbing	A	64	-0232		
10.	Hydrogen Drying (Molecular Sieves)	В	32	.0117		
11.	Crude TCS Recycle System	В	58	.0212		
12.	Silicon Fines Sep- aration	B	15	.0055		
13.	Material Handling	A			90	.0329
14.	Polysilicon Production		732	.2672		
	TOTAL		1386	.5059	90	.0329

NOTES:

^{1.} A Batch Process or Multiple Small Units

B Average Process

C Automated Process

^{2.} Man hours/day Unit from Figure 4-6, Peters and Timmerhaus (7).

^{3.} Polysilicon manpower requirements based on batch operation with approximately 1 operator per 10 reactors.

^{4.} Filament puller manpower requirements based on 1 operator per puller.

3.3 UCC Silane Process for Silicon (Union Carbide Corporation)

The chemical engineering analysis activity involves a preliminary process design of a plant to produce silicon via the technology under consideration.

The UCC silane process (Union Carbide Corporation) for silicon involves several processing operations of hydrogenation-hydrochlorination reaction, stripping, distillation, redistribution reaction, silane purification, pryolysis and consolidation of silicon. The process flowsheet is shown in Figure 3.3-1.

Hydrogen, silicon tetrachloride, and metallurigical grade silicon are fed to the hydrogenation reactor (fluidized bed, 500°C, 515 psia, copper catalyst) to produce a mixture of chlorosilanes. The mixture of chlorosilanes from the hydrogenation reaction is condensed and subjected to several state distillation to separate components and remove impurities.

Initially, the condensed liquid mixture is sent to D-01 stripper (90 psia) to remove inert gases and volatile impurities. The stripper bottoms go to D-02 distillation (55 psia) which separates TCS (trichlorosilane) and STC (silicon tetrachloride). The TCS redistribution reactor (liquid phase, 85 psia, 140°F catalyst) is used to produce DCS (dichlorosilane). The separation of DCS and TCS is achieved in D-03 distillation (320 psia). The overhead goes to DCS redistribution reactor (liquid phase, 510 psia, 140°F, catalyst) to produce silane (SiH₄). The silane is purified by separation from trace impurities (such as $B_{\rm pH_G}$) by D-04 distillation (355 psia).

The purified situae is used to produce silicon powder via the pyrolysis reaction:

$$SiH_4 + Si + 2H_2$$
. (3.3-1)

The hydrogen from the reaction is compressed and recycled to the hydrogenation reactor. The silicon powder from the pyrolysis is consolidated to provide the molten silicon product.

A process design was performed to obtain data for a cost analysis of a plant to produce silicon by this new technology. The design was based on a plant to produce 1000 metric tons/yr of silicon via the UCC silane process.

The detailed status sheet for the process design package is shown in Table 3.3-1, and is representative of the various sub-items that make up the activity. The summarized results for the preliminary process design are presented in a tabular format to make it easier to locate items of specific interest.

The guide for these tables is given below:

```
•Process Flowsheet------Figure 3.3-2
•Base Case Conditions------Table 3.3-2
•Reaction Chemistry------Table 3.3-3
•Redistribution Equilibrium-----Figure 3.3-3
•Raw Material Requirements-----Table 3.3-4
•Utility Requirements-----Table 3.3-5
•Major Process Equipment------Table 3.3-6
•Production Labor Requirements----Table 3.3-7
```

The process design provides detailed data for raw materials, utilities, major process equipment and production labor requirements which are necessary for polysilicon production.

Figure 3.3-1 PROCESS FLOWSHEET FOR UCC SILANE PROCESS

TABLE 3.3-1 CHEMICAL ENGINEERING ANALYSES:
PRELIMINARY PROCESS DESIGN ACTIVITIES FOR UCC SILAME PROCESS

	Prel. Process Design Activity	Status		Prel. Process Design Activity	Status
1.	Specify Base Case Conditions 1. Plant Size	•	7.	Equipment Design Calculations 1. Storage Vessels	•
	2. Product Specifics3. Additional Conditions	;		 Unit Operations Equipment Process Data (P, T, rate, etc.) Additional 	•
2.	Define Reaction Chemistry	•			
	 Reactants, Products 	•	8.	List of Major Process Equipment	•
	2. Equilibrium	•		l. Size	•
				2. Type	•
3.	Process Flow Diagram	•		3. Materials of Construct:on	•
	1. Flow Sequence, Unit Operations	•			
	<pre>1. Process Conditions (T, P, etc.)</pre>	•	8a.	Major Technical Factors	•
22	s. Environmental	•		(Potential Problem Areas)	•
ယ	4. Company Interaction	•		l. Materials Compatibility	•
	(Technology Exchange)			 Process Conditions Limitations Additional 	•
4.	Material Balance Calculations	ŧ			
	1. Raw Materials	9	9.	Production Labor Requirements	•
	2. Products	•		1. Process Technology	•
	3. By-Products	•		2. Production Volume	•
5.	Energy Balance Calculations	•	10.	Forward for Economic Analysis	•
	1. Heating	•			
	2. Cooling	•			
	3. Additional	•			
6.	Property Data	•		0 Plan	
	1. Physical	•			
	2. Thermodynamic	•		● Complete	
	Additional	•			

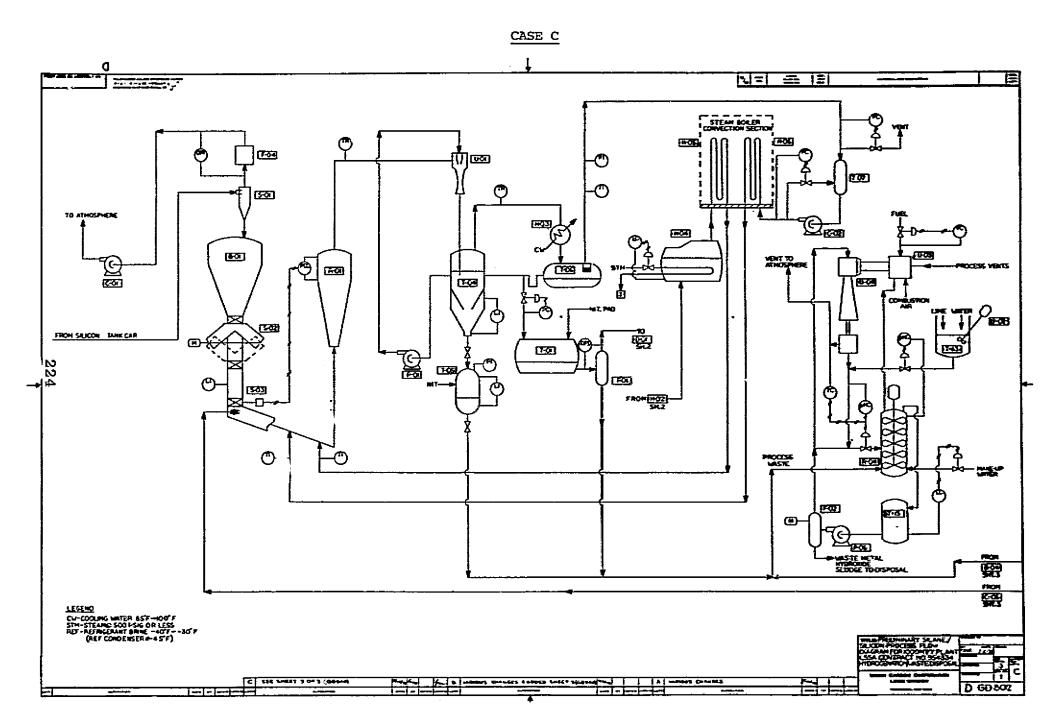


Figure 3.3-2 Process Flow Sheet for UCC Silane Process

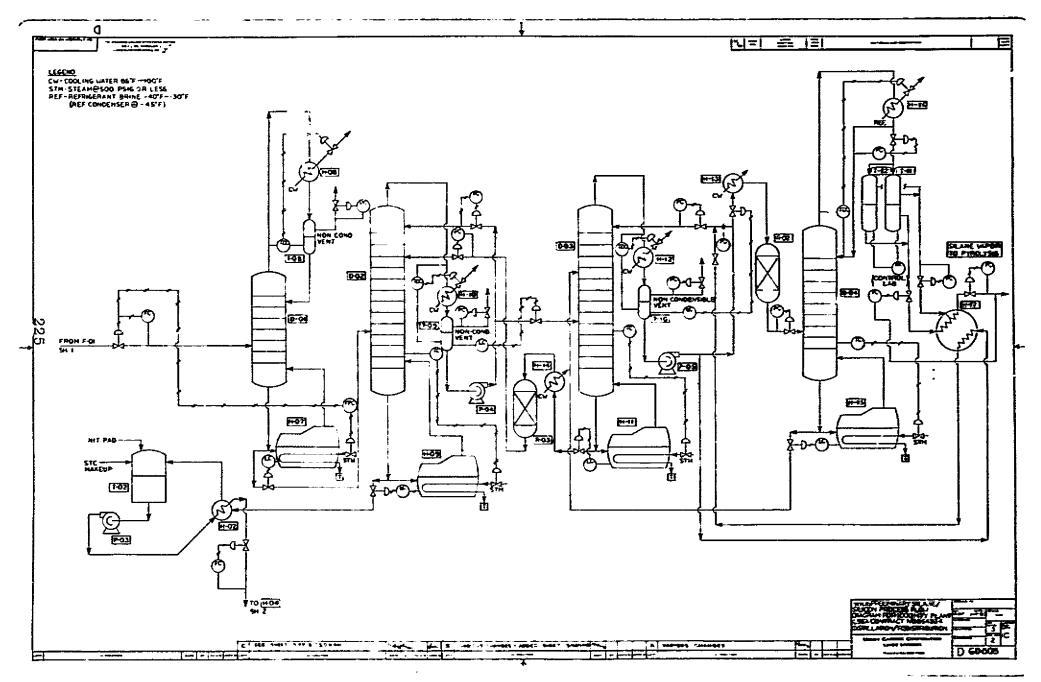


Figure 3.3-2 (Continued)

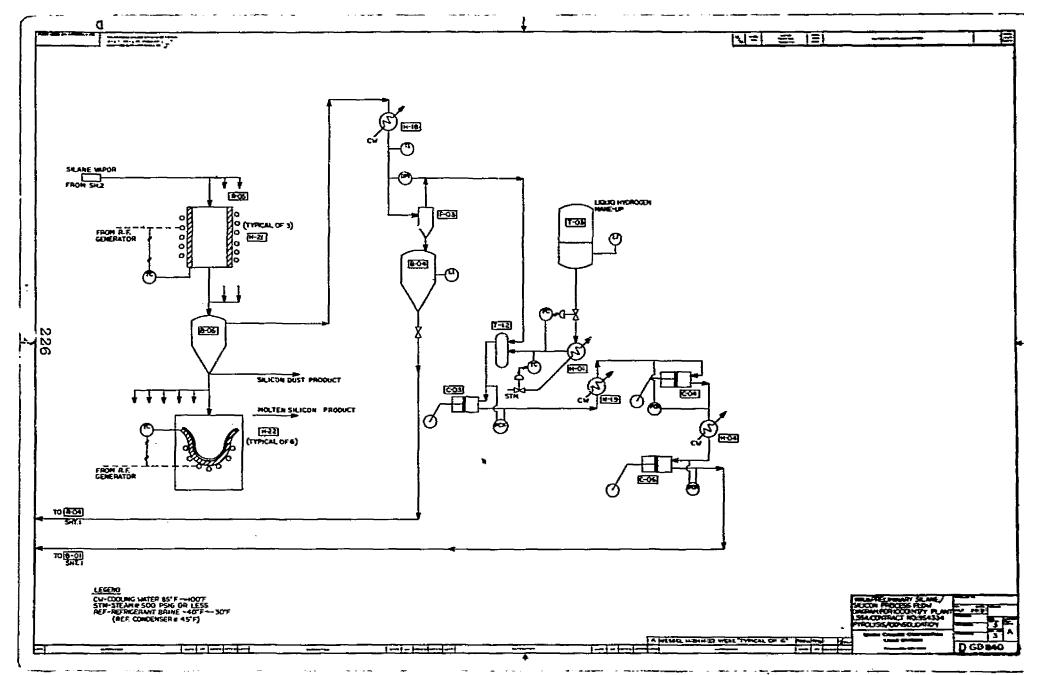


Figure 3.3-2 (Continued)

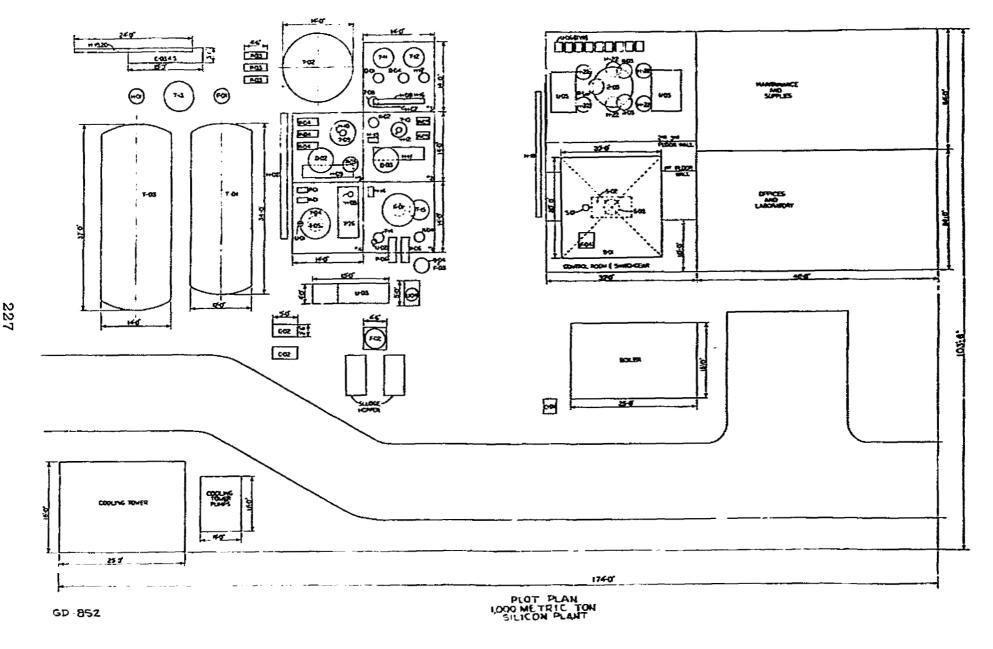


Figure 3.3-2 (Continued)

TABLE 3.3-2

BASE CASE CONDITIONS FOR UCC SILANE PROCESS

- 1. Plant Size
 - -Silicon produced from silane
 - -1000 metric tons/year of silicon
 - -Bolar cell grade silicon
- 2. Hydrogenation Reaction
 - -Metallurgical grade silicon, hydrogen, and recycle silicon tetrachloride (TET) used to produce trichlorosilane (TCS)
 - -Copper catalyzed
 - -Fluidized bed
 - -500°C, 514.7 psia
 - -20% to 22.5% conversion of SiCl4 (example)
- 3. TCS Redistribution Reaction
 - -TCS from hydrogenation produces dichlorosilane (DCS)
 - -Catalytic redistribution of TCS with tertiary amine ion exchange resin
 - -Liquid phase 85 psia, 140°F
 - -Conversion a function of intet concentration (Union Carbide equilibrium)
 - -Conversion from pure TCS feed as about 9.5% to DCS (example)
- 4. DCS Redistribution Reaction
 - -DCS produces SiH4 (silane)
 - -Catalytic redistribution of DCS with tertiary amine exchange resin
 - -Liquid phase 510 psia, 140°F
 - -Conversion a function of inlet concentration (Union Carbide equilibrium)
 - -Conversion from pure DCS feed is about 14% to Silane (example)
- 5. Recycles
 - -Unreacted chlorosilanes separated by distillation and recycled
- 6. Silane Purification
 - -Final purification by distillation
 - -Designed to remove trace impurities (B2H6, example)
- 7. Operating Ratio
 - -Approximately 85% utilization (on stream time)
 - -Approximately 7445 hour/year production
- 8. Storage Consideration
 - -Feed materials (several week supply, approx. 1 month)
 - -Product (two shifts storage)
 - -Process (several hours to 1 shift)

TABLE 3.3-3

REACTION CHEMISTRY FOR UCC SILANE PROCESS

1. Hydrogenation Reaction

$$3 \operatorname{sicl}_4 + \operatorname{si} + 2\operatorname{H}_2 \div 4 \operatorname{siHCl}_3$$

2. Trichlorosilane Redistribution Reaction

$$2 \operatorname{siHCl}_{3} \stackrel{?}{+} \operatorname{SiH}_{2}\operatorname{Cl}_{2} + \operatorname{siCl}_{4}$$

3. Dichlorosilane Redistribution Reaction

4. Waste Treatment (representative)

5. Silane Pyrolysis Reaction

$$siH_4 \longrightarrow si + 2H_2$$

Note

- Reaction 1 product contains H₂, HCl, SiCl₄, SiHCl₃, SiH₂Cl₂(trace), other trace chlorides
- 2. Reaction 2 Product contains SiHCl₃ SiCl₄, SiH₂Cl₂, SiH₃Cl
- 3. Reaction 3 Product contains SiH_2Cl_2 , $SiHCl_3$, $SiCl_4$, SiH_3Cl , SiH_4

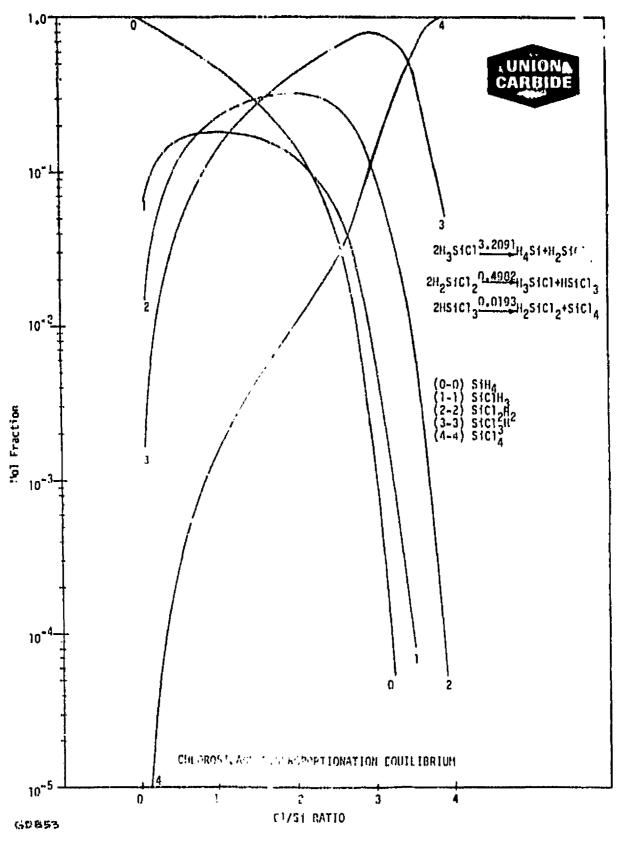


Figure 3.3-3 Red.strib dien Equilibrium For UCC Silane Process (Provided by Onton Carbide)

TABLE 3.3-4

RAW MATERIAL REQUIREMENTS FOR UCC SILANE PROCESS

	Raw Material	Requirements			
		lb/hr for 1000 MT/yr Silicon	lb/KG of Silicon		
1.	M. G. Silicon (Si)	348.6	2.60		
2.	Silicon Tetrachloride (SiCl ₄ , make-up)	370.1	2.76		
3.	Liquid Hydrogen (H ₂ , make-up)	4.3	0.032		
4.	Copper Catalyst (Cu)	6.8	0.051		
5.	Hydrate Lime [Ca(OH)2]	326.3	2,43		

Note:

Assuming all inert gas from D-O1 is ${\rm H_2}$

TABLE 3.3-5
UTILITY REQUIREMENTS FOR UCC SILANE PROCESS

Utility/Function	Total Requirement	Requirement/KG Silicon
l. Electricity	409.7 KW	3.050 KW/hr
2. Steam	23113 lb/hr	172.2 lbs.
3. Cooling Water	70,453 gal/hr	525 gallons
4. Process Water	9.52 gal/hr	0.0709 gallons
5. Refrigerant	0.130 MM Btu/hr	968 Btu
6. Fuel	3.640 MM Btu/hr	27,100 Btu

TABLE 3.3-6

LIST OF MAJOR PROCESS EQUPIMENT FOR UCC SILAME PROCESS

		Equipment	Function	Duty/Type	<u>Size</u>	Material of Construction
	DIS	TILLATION COLUMNS				
	1.	D-01 Crude TCS/STC Stripping Column	To remove inert gases	24929 lb/hr of feed	24" diam. 20' tall, 10 sieve plates	cs
	2.	D-02 TCS/STC Distillation Column	To remove STC at bottoms	64,213 lb/hr of feeds	4.63' diam., 74' tall, 32 sieve plates	cs
	3.	D-03 DCS/TCS Distillation Column	To remove DCS at distillates	46,254 lb/hr of feeds	4.30' diam., 77.5' tall, 45 sieve plates	cs
233	4.	D-04 Silane Distil- lation Column	To purify silane	6967 lb/hr of feed	24" diam., 50'tall, 30 sieve plates	316SS
	REA	ACTORS				
	5.	R-Ol Hydrogenation Reactor (Fluidized Bed)	Hydrogenation of Si and SiCl ₄	25,447 lb/hr of feed	6.5' diam. x 6.5'/ 2.5' diam. x 10.5' cone	316SS
	6.	R-02 DCS Redistri- bution Reactor (Fixed Bed)	To convert DCS to silane	6,967 lb/hr of feed	2' diam., 11.2' tall, with catalyst	316SS
	٦.	R-03 TCS Redistrı- bution Teactor (Fixed Bed)	To convert TCS to DCS and STC	39,287 lb/hr of feed	3' diam., 17.3' tall, with catalyst	31 6 SS
8. 8	§ 9.	R-04 Sludge Neutralization Reactor	Waste Treatment	Agitated Tank/Colvæn	2' diam., 20' tall	316ss (for pricing)

HEAT EXCHANGERS

	10.	H-Ol Liquid Hydrogen Vaporizer	To provide H ₂ gas	7.22 lb/hr of Liquid H ₂	(Vendor supplied equipment)	
	11.	H-O2 STC Cooler	Exchange heats of STC streams	7.23 x 10 ⁵ Btu/hr shell-tube H. E.	893 ft. ² 514.7 psia	316SS/CS
	12.	H-03 Quench Condenser	To condense chlorosilanes, 100°F	4.69 x 10 ⁶ Btu/hr shell-tube H. E.	676 ft. ² 514.7 psia	316SS/CS
234	13.	H-04 Recycle STC Vaporizer	To provide STC vapor to reactor	l.71 x 10 ⁶ Btu/hr kettle	65.9 ft. ² 514.7 psia	CS
44	14.	H-05 Recycle STC Superheater	To heat STC from 234 to 932°F	2.46 x 10 ⁶ Btu/hr Furnace Convection	1603 ft. ² 514.7 psia	316SS
	15.	H-06 Recycle H ₂ Heater	To heat H ₂ from 100 to 932°F	6.78 x 10 ⁵ Btu/hr Furnace Convection	331 ft. ² 514.7 psia	316SS
	16.	H-07 Stipper Recoiler	Reboiler of 9-01, 242°F	9.06 x 10 ⁵ Btu/hr Kettle	39.8 ft. ² 95 psia	cs
	17.	H-08 Stripper Condenser	Partial Condenser of D-Ol, 139°F	86,700 stu/hr shel -tu e 8 %.	36.1 ft. ² 90 psia	cs
	18.	H-09 TCS/STC Reboiler	Reboiler of D-O2 216°F	7.83 x 10° Bcu/hr kettle	295 ft. ² 55 psia	cs
	19.	H-10 TCS/STC Condenser	Total condenser of D-02, 120°F	6.24 x lo' Btu/hr shell-tube H. E.	1315 fc. ² 55 psia	316SS/CS

20.	H-11 DCS/TCS Reboiler	Reboiler of D-03	3.50 x 10 ⁶ Btu/hr Kettle	333 ft. ² 320 psia	CS
21.	H-12 DCS/TCS Condenser	Total condenser of D-03, 234°F	3.99 x 106 Btu/hr shell-tube H. E.	429 ft. ² 320 psia	316SS/CS
22.	H-13 DCS Cooler	To cool DCS before redistribution reaction	1.88 x 10 ⁵ Btu/hr shell-tube H. E.	22.1 ft. ² 550 psia	316SS/CS
23.	H-14 TCS Cooler	To cool TCS before redistribution reaction	2.01 x 10 ⁶ Btu/hr shell-tube H. E.	161 ft. ² 85 psia	316SS/CS
24.	H-15 Silane Boiler	Reboiler of D-04 278°F	2.71 x 10 ⁵ Btu/hr Kettle	15.4 ft. ² 360 psia	316SS/CS
25.	H-16 Silane Condenser	Total conderser of -44PF	1.30 x 10 ⁵ Btu/hr Shell-tube H. E.	81 ft. ² 360 psia	316SS
26.	H-17 Silane Vaporizer/Superheate:	To provide silane vapor for pyroly- sis, 200°F	25,000 Btu/hr Jacket/tubes	27.3 ft. ³ /12.0 ft. ² 355 psia	316SS/CS
27.	H-18 Pyrolysis Hydrogen Cooler	To cool H ₂ gas from 363 to 100°F		80.5 ft. ² 20 psia	3165S/CS
28.	H-19 First Stage H ₂ Intercooler	To cool H ₂ be- tween comp. stages 328 to 100°F	38,570 Btu/hr shell-tube H.E.	72.2 ft. ² 50 psia	316SS/CS
29.	H-20 Second Stage H ₂ Intercooler	To cool H ₂ be- tween comp. stages 328 to 100°F	38570 Btu/hr shell-tube H. E.	72.2 ft. ² 160 psia	31655/CS

PUMP	S AND COMPRESSORS				
30.	C-01 Pneumatic Conveying Fan	Si feed transport	417 ACFM Centrifugal	5.1 psi Δp 12 BHP	CS
31.	C-02 Recycle H ₂ Blower	H ₂ gas blower	22.8 ACFM Centrifugal	518.7 psia 1.23 BHP	CS
32.	C-03 First Stage H ₂ Compressor	H ₂ gas compressor	164 ACFM; double action, reciprocating	48.1 psia discharge, 19.8 BHP	cs
33.	C-04 Second Stage H ₂ Compressor	H ₂ gas compressor action,	50 ACFM; double action, reciprocating	157 psia discharge, 19.6 BHP	cs
34.	C-05 Third Stage H ₂ Compressor	H ₂ gas compressor	15.3 ACFM;double action, reciprocating	515 psia discharge, 19.8 BHP	CS
35.	P-01 Quench Contactor Pump	Circulating liquid chlorosilanes	100 gpm, centrifugal/motor	36.7' head, 1.56 BHP	316SS
36.	P-03 Recycle STC Ритр	To supply STC	33.1 gpm Centrifugal/ Turbine	847' head, 14.2 BHP	CS
37.	P-04 TCS Distillate Pump	D-02 Reflux/ Distillate	144 gpm Centrifugal/ Turbine	589' head, 33.5 BHP	CS
38.	P-05 DCS Distillate Pump	D-03 Reflux/ Distillate	144 gpm Centrifugal/ Motor	759' head, 26.2 BHP	316ss
39.	P-06 Lime Tank Pump	Circulating Lime Slurry	100 gpm Centrifugal/	103' head, 4.BHP	Cast iron

Motor

TANKS AND BINS

40.	T-01 Crude TCS/STC Storage Tank	Storage/Feed to Silane production	8 hr. storage, horizontal	12' diam. x 27' >sia	CS
41.	T-02 STC Storage Tank	Storage/Feed to Hydrogenation	6 hr. storage, Vertical	14' diam. x 13.4' 14.7 psia	CS
42.	T-03 Liq. H ₂ Storage	Liq. H ₂ make-up storage	(Vendor supplied eq	uipment)	
43.	T-04 Waste Settler Tank	To separate solid residues	Vertical cyl/cone bottom	6' diam., 12' tall 514.7 psia	316SS
44.	T-05 Waste Chlorides Tank	To remove solid residues	285 lb/hr Vertical	3' diam., 4' tall 25 psia (aprox.)	316SS
45.	T-06 Quench Condenser Receiver	Gas-liq. separa-	l ^r min. storage Horizontal	4' diam. x 11.3' 514.7 psia	cs
46.	T-07 Recycle Hydrogen Receiver	H ₂ gas surge tank	Vertical	3' diam. x 6' 514.7 psia	cs
47.	T-08 Stripper Reflux pot	D-01 Distillate/ gas	30 min. storage Vertical	2' diam. x 3.4' 90 psia	CS
48.	T-09 TCS/STC Reflux pot	D-02 Distillate	10 min. storage Vertical	5' diam. x 10' 55 psia	CS
49.	T-10 DCS/TCS Relux pot	D-03 Distillate	10 min. storage Vertical	4' diam. x 12' 320 psia	cs
50.	T-11 A, B Silane Shift Tanks	D-04 Distillate/ Feed to pyrolysis	4 hr. storage, each, Vertical	5' diam. x 9.4' 360 psia	316SS

TABLE	3	3-6	(Continued)
THEFT	- 7	36	(CONTINUED)

	51.	T-13 Pyrolysis H ₂ Receiver	H ₂ Feed to Com- pressor	Vertical 25 psia	6' diam. x 12'	cs
	52.	T-14 Lime Make-Up Tank	Lime solu. pre- paration	8 hr. storage Vertical, open	5' diam. x 9.2' 14.7 psia	CS
	53.	T-15 Sludge Pump Tank	Sludge-solu. storage	4 hr. storage Vertical	5' diam. x 8'	316SS
	54.	B-01 M. G. Silicon Storage Hopper	Feed to hydro- genation reactor	344 lb/hr Si	20' sq. x 5'/10' cone 14.7 psia	CS
55. &	56•	B-04 Pyrolysis Dust Bin	Solid residue	Small	3'diam. x 3' 25 psia	cs
238						
	FILT	TERS				
	57.	F-01 Crude TCS/ STC Filter	In Line Filter	Small solic-liq.	50 gpm x 100µ	CS/316SS
	58.	F-02 Waste Hydroxide Filter	Remove solid residues	100 gpm solid-liq.	Filtering area 67 ft. ²	CS/316SS
	59.	F-03 Pyrolysis H ₂ Filter	Remove solid residues	Small Bag solid-gas	20 ft. ² x 5µ	CS/cloth
	60.	F-04 M. G. Silicon Unloading Filter	Solid-air separa- tion in pneumatic conveyor		20 ft. ² x 5μ	CS/cloth

Ŋ

TABLE 3.3-6 (Continued)

SOLI	D HANDLING EQUIPMENT					
61.	S-01 M. G. Silicon Unloading Cyclone	Si feed trans- port	6" W. C. ΔΡ	417 ACFM	316SS	
62.	S-02 Double Shell Blender	Si feed to reactor	Blending	20 ft. 3/shroud	304SS	
63.	S-03 M. G. Silicon Lock Hopper	Si Feed to reactor	Locking	20 ft. ³	cs	
UNCL	ASSED					
64.	U-01 Quench Contactor Ejector	To withdraw gaseous products from reactor	6" W. C. Suction	100gpm 134 ACFM	31655	
65.	U-02 Lime Tank Agitator	Line solution preparation		3/4 HP	316SS	
66.	U-03 Vent Gas Combustor	To burn vent gases from various mits	_	1.22 ACFM	316SS	
67.	U-04 Vent Gas Sjector	To withdraw gases from combustion chamber	10" W. C. Suction	100gpm 1.0 CFM (STP)	316SS	
PYROLYSIS SECTION (Primary)						
68.	R-05 Silane Pyrolysis Reactors (six)	To convert silane to sil.con	25 KW Power supply	3' dram., 15' tall, cone shape (approx.)	Monel/ _uartz (pricing)	
69.	X-01 Melters (six)	Melt silicon	60 KW Power supply		Graphite/ Quartz, etc.	

70.	B-05 Powder Hoppers (six)	Pyrolysis Powder collection	96.1 lb/hr Si	8' diam. x 15' width cone bottom	316SS
71.	X-02 Hydrogen Cooler	Cool Hydrogen	42.4 lb/hr H ₂	See Economic Analysis	CS/Other
72.	X-03 Hydrogen Blower	Blow Hydrogen	120 CFM (STP)	See Economic Analysis	CS/Other
73.	X-04 Dust Filter	Filter Dust	127 CFM (STP)	See Economic Analysis	CS/Other
74.	X-05 Star Valve (six)	Flow Control		See Economic Analysis	CS/Other
75.	X-06 Conveyor	Transport Material	96.1 lb/hr Si and its containers	See Economic Analysis	CS/Other
76.	X-07 Drum Loader	Load Drums	TES CONCUMICES	See Economic Analysis	

TABLE 3.3-7
PRODUCTION LABOR REQUIREMENTS FOR UCC SILANE PROCESS

Section/Unit Operation		Shilled Labor man-hr/KG Si (oper/shift)	Semiskillud Labor man-hr/KG Si (oper/shift)	
1.	Hydrogenation	0.00745 (1)	0.00745 (1)	
2.	Silane	0.02230 (3)		
3.	Pyrolysis	0.02980 (4)	tor on po ye	
4.	Waste Treatment	0.00745 (1)	64 TH 107 TH	
5.	Hydrogen Compression	0.00745 (1)	they tree gast from	
	TOTAL,	0.0745 (10)	0.00745 (1)	

Note

Manpower estimate for production labor requirements based on:

- Dividing plant into sections

 type of unit operation
 mark off working area
- 2. Specify work duties required in each section
- 3. Estimate operators required to perform work duties in each section
 - -type of unit operation
 - -size of working area
 - -degree of automation (batch, semi-continuous, continuous, etc.)

3.4 BCL Process for Silicon - Case A (Battelle Columbus Laboratories)

The chemical engineering analysis activity involves a preliminary process design of a plant to produce silicon via the technology under consideration.

The process flowsheet for the Case A of BCL process to manufacture silicon is shown in Figure 3.4-1. This process consists of several major processing operations of distillation, vaporization, stripping, condensation and a deposition reaction to produce silicon as well as electrolysis to recover the zinc.

Silicon tetrachloride (SiCl₄), which is the major raw material, is fed to the distillation section for purification, to remove impurities (such as boron and phosphorous). In the deposition section, purified silicon tetrachloride is vaporized and preheated to the reaction temperature, 927°C, before it is introduced into a silicon deposition unit, which is a fluid-ized bed reactor. Zinc vapor, produced by a specially designed induction-heated vaporizer, is also introduced to the reactor at the same temperature for the reaction. The reaction equation to show the silicon deposition is

$$SiCl_4 + 2Zn + Si + 2ZnCl_2$$
 (3.4-1)

Silicon granules produced by the deposition reaction, which descend to the bottom of reactor, are cooled and collected in containers. A small amount of silicon seed is fed to the reactor to control the particle size of the silicon product. Zinc chloride and unreacted zinc are recovered and fed to the electrolysis section, while unreacted silicon tetrachloride is recycled to the distillation section.

In the electrolysis section, zinc chloride is reduced to zinc by low voltage (4-5 volts) electrolysis cells. Zinc is recycled to the deposition unit, while chlorine gas is collected as the by-product. The deposition and electrolysis sections are purged with inert can (such as argon). Waste gases from various sections are collected and treated with hydrate lime solution in the waste treatment section.

A process design was performed to obtain data for a cost analysis of a plant to produce silicon by this new technology. The design was based on a plant to produce 1000 metric tons/yr of silicon via the BCL process. In Case A, two deposition reactors and six electrolysis cells are required.

The detailed status sheet for the process design package is shown in Table 3.4-1 and is representative of the various sub-items that make up the activity. The summarized results

for the preliminary process design are presented in a tabular format to make it easier to locate items of specific interest. The guide for these tables is given below:

The process design provides detailed data for raw materials, utilities, major process equipment and production labor requirements which are necessary for polysilicon production.

TABLE 3.4-1 CHEMICAL ENGINEERING ANALYSES:
PRELIMINARY PROCESS DESIGN ACTIVITIES FOR BCL PROCESS - Case A

	Prel. Process Design Activity	<u>Status</u>	Prel. Process Design Activity Status
1.	Specify Base Case Conditions	•	7. Equipment Design Calculations
	1. Plant Size	•	1. Storage Vessels
	2. Product Specifics	•	2. Unit Operations Equipment
	Additional Conditions	•	3. Process Data (P, T, rate, etc.)
			4. Additional •
2.	Define Reaction Chemistry	•	
	 Reactants, Products 	•	8. List of Major Process Equipment
	2. Equilibrium	•	1. Size
			2. Type ●
3.	Process Flow Diagram	•	3. Materials of Construction
	1. Flow Sequence, Unit Operations	•	
	2. Process Conditions (T, P, etc.)	•	8a. Major Technical Factors
	 Environmental 	•	(Potential Problem Areas)
	4. Company Interaction	•	l. Materials Compatibility •
	(Technology Exchange)		2. Process Conditions Limitations
			3. Additional
4.	Material Balance Calculations	•	
	l. Raw Materials	•	9. Production Labor Requirements
	2. Products	•	1. Process Technology
	3. By-Products	•	2. Production Volume
5.	Energy Balance Calculations	•	10. Forward for Economic Analysis
	l. Heating	•	
	2. Cooling	•	
	3. Additional	•	
6.	Property Data	•	0 Flan
	1. Physical	•	0 In Progress
	2. Thermodynamic	•	● Complete
	3. Additional	•	

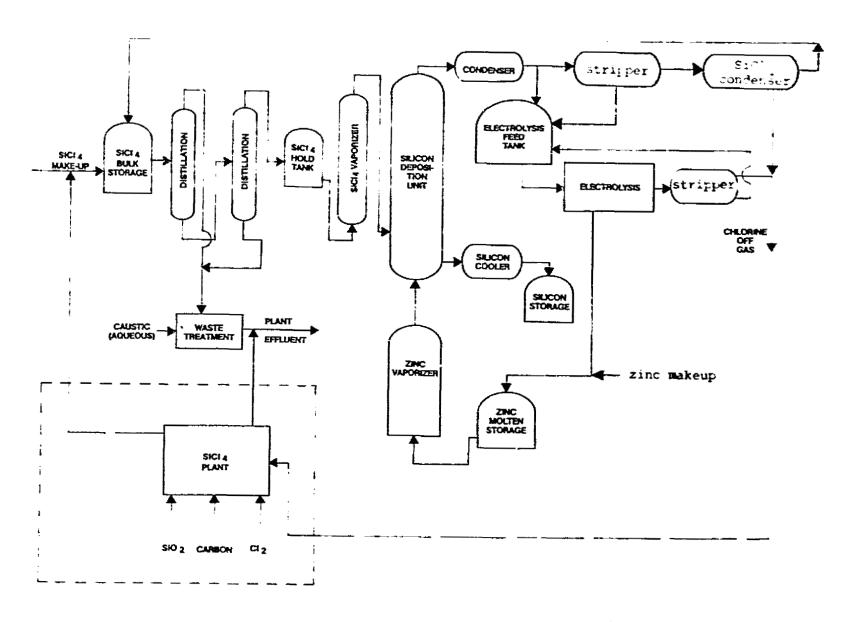


Figure 3.4-1 Process Flow Sheet for BCL Process-Case A

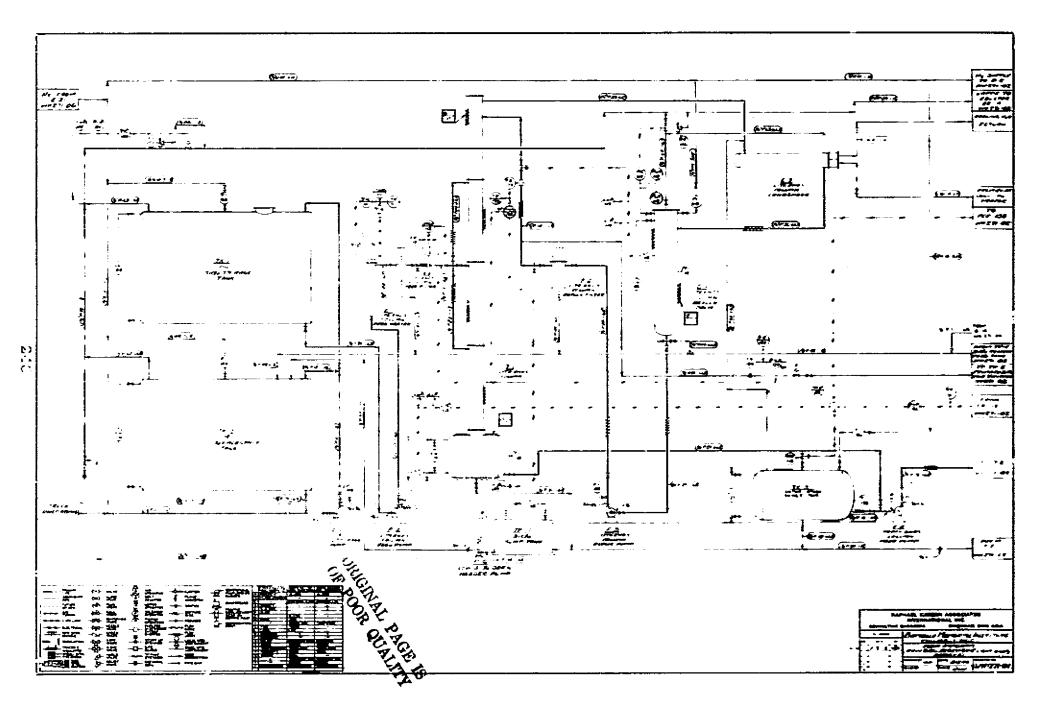


Figure 3.4-2 Process Flow Sheet for BCL Process-Case A

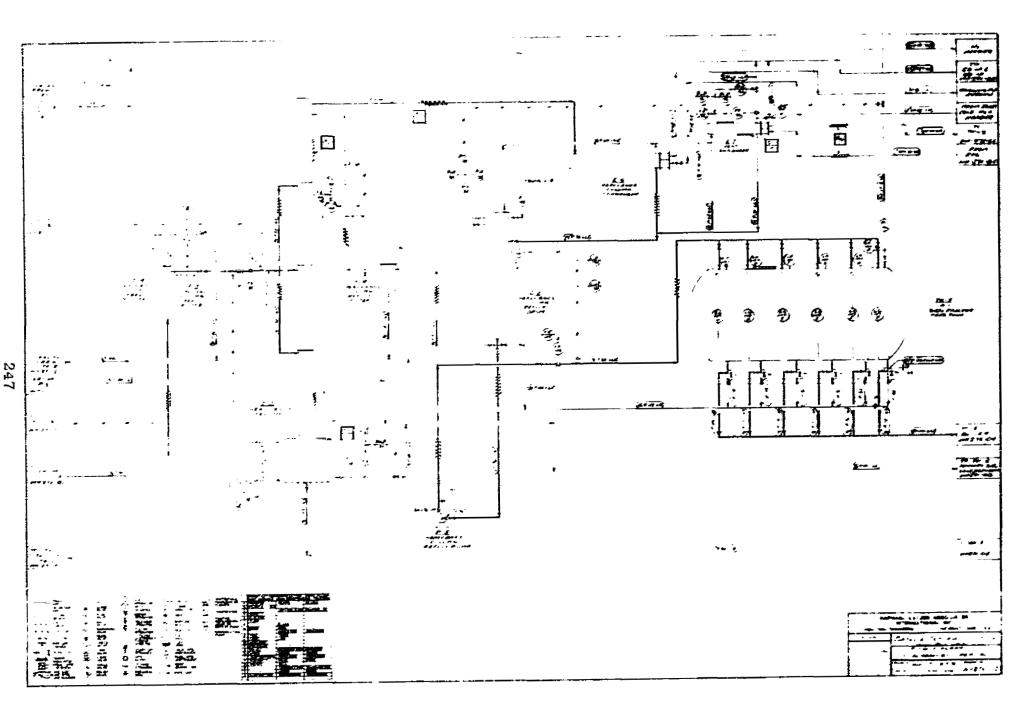


Figure 3.4-2 (Continued)

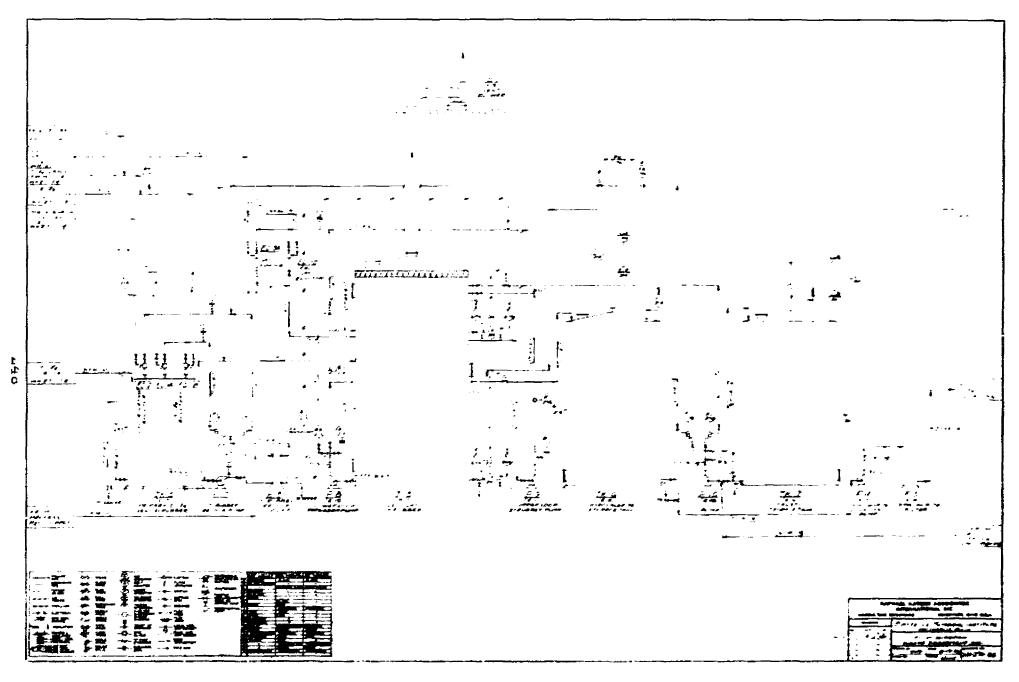


Figure 3.4-2 (Continued)

Note: For 50 MT/yr

Figure 3.4-2 (Continued)

Figure 3.4-2 (Continued)

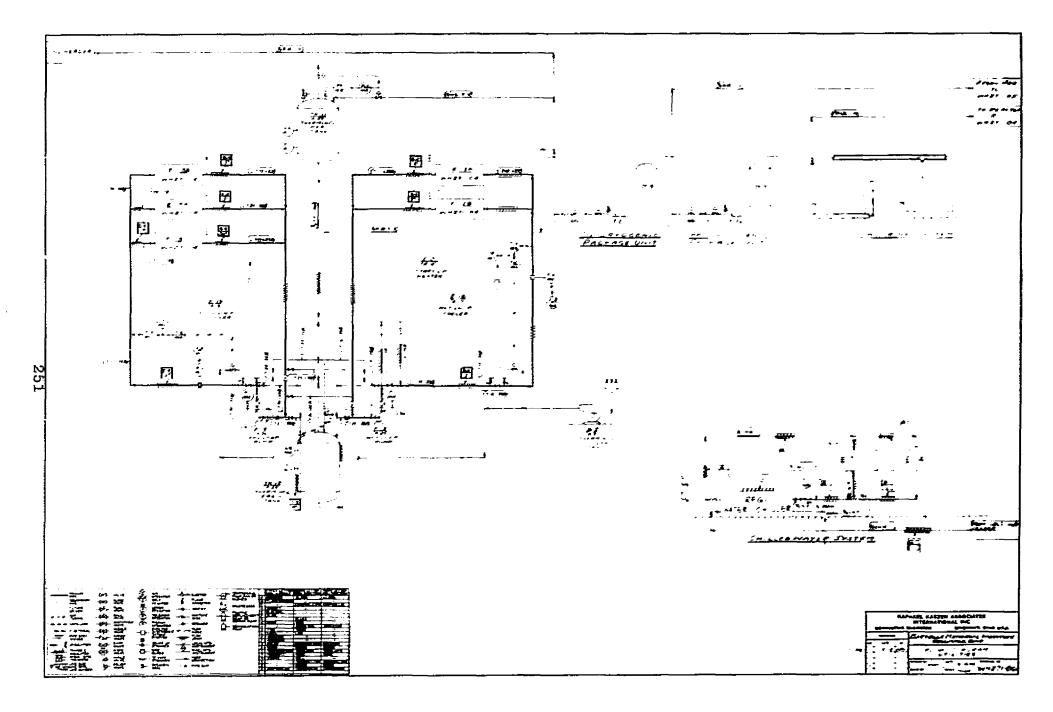
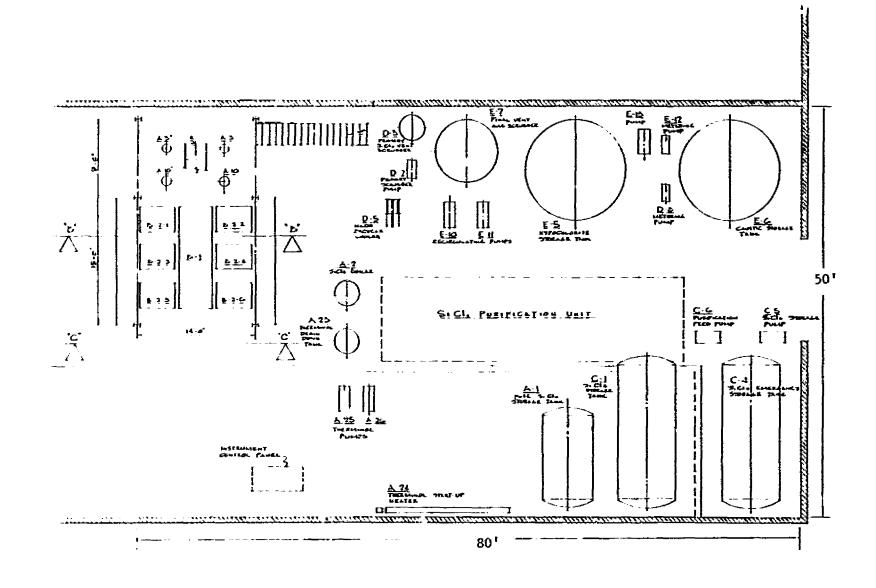


Figure 3.4-2 (Continued)



PLOT PLAN

Figure 3.4-2 (Continued)

TABLE 3.4-2 BASE CASE CONDITIONS FOR BCL PROCESS - Case A

- 1. Plant Size
 - -silicon produced from silicon tetrachloride (TET)
 - -1000 metric tons/yr of milicon
 - -solar cell grade silicon
 - -solid phase product form (granules)
- Light End Distillation
 - -purification of TET by distillation
 - -remove 4% chlorosilanes as the light end
 - -80°C, 10 paig
- 3. Heavy End Distillation
 - -purification of TET by distillation
 - -remove 4% impurities as the heavy end
 - -92% over-all yield of TET from both distillations
 - -80°C, 10 psig
- 4. TET Vaporizer
 - -to supply TET vapor for deposition reactor
 - -by power input (resistance heater)
 - -hold at constant level and constant pressure
 - -164°F
- 5. Deposition Reactor
 - -reduce TET by zinc to produce silicon
 - -deposit on pure silicon seed
 - -fluid bed
 - -927°C (1700°F, 1 atm)
 - -63% conversion of TET to silicon
- 6. Reactor Condenser
 - -to condense games from reactor (ZnCl₂, unreacted Zn and SiCl_A games)
 - -partial condensation
 - -using therminol 66 as the coolant
 - -927°C inlet temperature and 350°C outlet temperature
- 7. Reactor ZnCl Stripper
 - -work as partial condenser
 - -to condense ZnCl_2 gas from SiCl_4 gas
 - -operating at the temperature right above ZnCl₂ melting point (318°C), 350°C -using therminol 66 as the coolant
 - Cell ZnCl, Stripper
 - -operates as partial condenser

 - -to condense $\rm ZnCl_2$ gas from $\rm Cl_2$ and $\rm SiCl_4$ gases -operating at the temperature right above $\rm ZnCl_2$ melting point (318°C), 350°C
 - -using therminol 66 as the heat exchange medium

- 9. Reactor SiCl4 Condenser
 - -condense SiCl4 gas for recycle
 - -antifreeze as the coolant
 - -350°C inlet temperature, 20°F outlet temperature.
- 10. Electrolysis
 - -electrolytic recovery of Zn from ZnCl2
 - -Cl2 gas is by product
 - -95% Zn recovery
 - -500°C, approx. I atm
- 11. Zinc Vaporizer
 - -to vaporize Zinc
 - -by induction heating
 - -927°C, approx. 1 atm.
- 12. Wastes Treatment
 - -to scrub and neutralize ${\rm SiCl}_4$ and chlorosilane gases -caustic solution used to neutralize
- 13. Operating Ratio
 - -approximately 80% utilization (on struct time)
 - -approximately 7,000 hr/yr production
- 14. Storage Considerations
 - -feed material (two week supply)
 - -product (two shifts storage)
 - -process (several hours)

TABLE 3.4-3 REACTION CHEMISTRY FOR BCL PROCESS - Case A

1. Silicon Deposition

$$2Zn + SiCl_4 + Si + 2ZnCl_2$$

2. Electrolysis

$$ZnCl_2$$
 electroly. $Zn(1) + Cl_2(g)$

3. Waste Treatment

$$sicl_4 + 2H_2O \xrightarrow{aq} sio_2 + 4Hcl(aq)$$

$$NaOH(aq) + HCl(aq) \rightarrow NaCl(aq) + H_2O$$

or

$$Ca(OH)_2(aq) + 2HCl(aq) + CaCl_2(aq) + 2H_2O$$

3a. Waste Treatment (50 MT/yr unit)

$$Cl_2(g) + 2NaOH(aq) \rightarrow NaOCI(aq) + NaCI(aq) + H_2O$$

Table 3.4-4

Raw Material Requirements for BCL Process - Case A

	Raw Material	Requirements lb/KG of Silicon
1,	Silicon Tetrachloride, SiCl4	15.68 ¹
2.	Zinc, Zn	0.54
ż	Caustic (50%), NaOH(aq)	5.23
	or	
	Lime (99%), Ca(OH) ₂	2.852
4.	Argon	3.1 gcF3
5.	Nitrogen	7,6 scr3
6.	Chlorine, Cl ₂ (by-product)	11.12

- 1. Includes light wastes (4%), heavey wastes (4%) and additional losses (7%).
- 2. Includes neutralization of distillation section, deposition section, electrolysis section and chlorination losses.
- 3. Estimate from BCL

TABLE 3.4-5

UTILITY REQUIREMENTS FOR BCL
PROCESS - Case A

		Utility/Function		Requirements/Kg of Silicon Product
1.	1. Electricity			30.92 kw-hr
		Low Volatage D.C. for Electro- lysis	(20.51)	
	2.	Zinc Vaporizer Induction Heated	(5.62)	
		Preheat Section of Deposition	(1,39)	
	-•	Unit Induction Heated		
	4.	Electrolysis Feed Tank Heater	(0.24)	
	5.		(0.10)	
		SiCl ₄ Vaporizer	(0.53)	
		Pumps, Blowers	(2.53)	
	• •	tanger, beautiful	(2002)	
2.	Ste	eam (50 PSIA)		9.67 pounds
		#1 Purification Column Reboiler	(4.59)	4 = 5 · · · · ·
	2.	#2 Purification Column Reboiler	(4.30	
	-	Caustic Storage Heating	(0,29)	
		#1 Purification Column Preheater	-	
	_	••		
3.		ling Water		37.88 Gallons
		#1 Purification Column Condenser		
		#2 Purification Column Condenser	(15.88)	
		Purified Tet Cooler	(1.67)	
	4.	Chlorination Cooler (H-17)	(0,53)	
	5.	Cell Gas Cooler (H-18)	(2.86)	
4.	Pro	cess Water		24.20 Gallons
		Diluent for Waste Treatment	(24.20)	
	•			
5.	Re f	rigeration		2.38 k9tu
	1.		(1.28)	
	2.	• •	(1.10)	

Note: $k = kilo = 10^3$

TABLE 3.4-6

LIST OF MAJOR PROCESS EXPLIPMENT FOR BCL - Case A

		Equipment	Function	Size/Type	Material of Construction	Capacity Ratio to 1000 MT/yr
	PRO	CESS TOWER AND INTERN	ALS			
	1.	D-01 Light End Distillation Column	To purify SiCl ₄	8" dia. x 21', packed 13.5'	Column, CS/packing, SS	20
	2.	D-02 Heavy End Distillation Column	To purify SiCl4	8" dia. x 21', packed 13.5'	Column, CS/packing, SS	20
	3.	A-01 Primary SiCl ₄ Vent Scrubber	To scrub SiCl ₄ vent gas	3' dia. x 4'4" T/T, 225 gal/flat bottom	FRP	1
2 5 5 8	4.	A-02 Final SiCl ₄ Vent Scrubber	To scrub SiCl ₄ vent gas	7'6" dia. x 17'4" T/T/ 4 pp trays, Teflon dimister	FRP	1
	HEAT	F EXCHANGER				
	5.	H-01 L.E. Column Feed Heater	To preheat feed to D-01	2' dia. x 5', 15,013 Btu/hr / external heater	cs	20
	б.	H-02 L.E. Column Rehoiler	Reboiler of D-01	2' dia. x 3', 51,522 Btu/hr / external heater	cs	29
	7.	H-03 L.E. Column Condenser	Total condenser of D-01	47,430 Btu/hr/shell-tube H.E.	CS	25
		H-04 H.E. Column Feed Heater	To preheat ford to D-02	2' dia. x 5', 14,331 Btu/hr/external heater	cs	20

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	9.	H-05 H.E. Column Reboiler	Reboiler of D-02	2' dia. x 3', 56,641 Btu/hr/external heater	cs	20
	10.	H-06 H.E. Column Condenser	Total condenser of D-02	52,292 Btu/hr/shell- tube H.E.	CS	20
	11.	H-07 SiCl ₄ Vent Condenser	Condense SiCl ₄ from vent gas	38 ft ² , 18,000 Btu/ hr//ell-tube H.E.	cs	20
	12.	H-08 SiCl ₄ Vaporizer		2.75' dia. x 3' T/T. 13.648 Etu/hr/ resistance heater	cs	20
	13.	H-09 Reactor Condenser	To condense by products from reactor	14 " dia. x 6.4', 126,237.2 Btu/hr	Graphite W/SS shell	20
259	14.	H-10 Reactor ZnCl ₂ Stripper	To condense ZnCl ₂	12 ft ² , 2,652 Btu/hr/ shell-tube H.E., finned U-tube	316 SS	20
	15.	H-11 SiCl4 Condenser	To condense SiCl ₄ gas for recycle	6,401 Btu/hr (x 4.62 = 29.573 Btu/hr)	316 SS	20
	16.	H-12 Cell ZnCl. Stripper	To condense ZnCl ₂ vapor	9,841.4 Btu/hr/shell- tube, H.E. (x 0.32)	Inconel 600	20
	17.	H-13 Therminol Cooler (cold circuit)	To soul Thermino) 56	68 ft ² , 11,000 Btu/hr/ shell-tube d.E., 500 psia	cs	20
	18.	H-14 Therminol Cooler (hot circuit)	To cool Therminol	262 ft ² , 120,000 Btu/hr/ shell-tube H.E., 500 psid		20
	19.	H-15 Start-up Heater	Thermise start up heaves	98,950 Btu/hr/U-tube 15', resistance heater	cs	20
		H-16 Silicon Product Cooler (two)	To occl the Si product from reactor	5,735 Btu/hr	sic	20

	20a.	H-17 Chlorination Cooler		20,000 Btu/hr, Area 200 ft ²	SS	1
	20b.	H-18 Cell Gas Cooler		1.08 x 10 ⁵ Btu/hr, Area 1805 ft ²	CS	1
	PROCI	ess and storage vessei	<u>re</u>			
	21.	T-01 SiCl ₄ Storage Tank	Storage/feed to purification	7' dia. x 16' T/T/ 4,600 gal	cs	20
	22.	T-02 SiCl ₄ Emergency Storage Tank	Storage/feed to purification	7' dia. x 1 ^{<'} T/T/ 4,600 gal	CS	20
	23.	T-03 L.E. Column Reflux Drum	To hold distillate for reflux	12" dia. x 4'/23 gal	CS	20
200	24.	T-04 Surge Tank	Surge Tank for D-01 bo'tom	3' dia. x 4'/200 gal	CS	20
	25.	T-05 Sump Tank	Sump for purification unit	3' dia. x 4'/200 gal	CS	20
	26.	T-06 H.E. Column Reflux Drum	To hold distillate for reflux	12" dia. x 4'/23 gal	CS	20
	27.	T-07 Pure SiCl ₄ Storage Tank	Storage feed to SiCl ₄ Vaporizer	6' dia. x 10' T/T/ 1900 ga¹	cs	20
	28.	T-08 Electrolysis Feed Tank	Storage feed ZnCl ₂ to electrolysis cel		Graphite/304 SS	20
	29.	T-09 Molter Zinc Storage Tank	Storage feed to Zinc vaporizer	W/heater 68,242 Btu/hr	Graphite/304 SS	20
	30.	T-10 Therminol Head Tank	Storage Therminol	1.5' dia. x 3.75' T/T/ 49.6 gal	CS	20

	31.	f-11 Therminol Drain Down Tank	To store drained Therminol	2.75' dia. x 3' T/T/ 133 gal	CS	20
	32.	T-12 Chlorine Sup- ply Tank	To supply chlorine gas	l 1/2' dia. x 3'/ 37.62 gal	cs	20
	33.	T-13 Lime Storage Tank	Storage Lime	12' dia. x 14'6" T/T/ 12,000 gal	PRP	1
	PUMP	S WITH DRIVERS				
	34.	P-01 Purification Feed Pump	To feed SiCl ₄ to storage tank	30 gpm, 31' head/ centrifugal, 1 1/2 hp	CS	20
	35.	P-02 L.E. Column Feed Pump	To supply SiCl ₄ to preheater	28.9 gph, Δ _F = 72 psia/ 0 ₋ 5 hp.	Cs	20
∾	36.	P-03 L.E. Column Relux Pump	D-01 Reflux	51.7 gph, $\Delta p = 23 \text{ psia/} 0.5 \text{ hp.}$	CS	20
61	37.	P-04 Surge Tank Pump	To supply SiCl ₄ to H.E. Column	29.4 gph, Δp = 53 psia/ 0.5 hp.	Cs	20
	38.	P-05 Sump Pump	To pump Sicl ₄ to emergency tank	30 gpm, 31' head/ centrifugal, 1 1/2 hp.	CS	20
	39.	P-06 L.E. Column Bottom Pump	To pump SiCI_4 to surge tank	29.4 gph, Lp = 53 psia/ 0.5 hp.	cs	20
	40.	P-07 H.E. Column Relux Pump	D-02 Reflux	57.1 gph, iv 25 psia/ 0.5 hp.	CS	20
	41.	P-98 H.E. Column: Bottom Pump	To pump bottom solution to waste treatment	1.3 gph, $\dot{c}_{r} \approx 25 \text{ psia/}$ 0.5 hp.	cs	2 0
	42	P-09 SiCl ₄ Vaporizer Feed Pump	To feed SiCl ₄ to Vaporizer	15 gph, 31' head/ 1/2 hp	CS	20

43.	P-10 Reactor Cin- denser Circulating Pump	To direulate con- densates	2.4 gpm, 30' head/1/2 hp	Graphite	20
44.	P-11 Cold Direct Pump (two)	Cold Therminol circulation	20 gpm, 85' head/centri- fugal, 2 hp.	CS	20
45.	P-12 Hot Circuit Pump	Hot Therminol circulation	62 gpm, 85' head/centri- fugal, 4 hp.	CS	20
45.	P-13 Primary Scrubber Redirbula- tion Pump	Recirculation for Scrubber A-01	20 gpm, 125' head/ centrigugal, 2.5 hp.	Duriron	1
; "	P-14 Primary Scrubber Lower-loop Recirculating Pump		100 gpm, 103' head/ centrifugal, 7 1/2 hp.	Duriron	1
48	P-15 Primary Scrubber Upper-loop Recirculating Pump	Circulate solution for upper-loop of Scrubber A-02	100 gpm, 13' head/ centrifugal, 2 hp.	Duriron	1.
49.	P-16 Make up Lime Metering Pump	Wime make up	0.9 gpm, 25' head/ centrifugal, 1/2 hp.	cs	1
FILT	ERS				
50.	F-01 L.E. Column. Feed Filter	Remove solids	29 gph, Δp = 5 psia/ 140 micron	Cs	20
51.	F-02 L.E. Column Reflux Filter	Remove solids	30 gph, Δp = 5 psia/ 140 micron	CS	20
52.	F-03 H. E. Column Feed Filter	Remove solids	52 gph, Δp = 5 psia/ 140 micron	CS	20
53.	F-04 H.E. Column Relfux Filter	Remove solids	31 gph, Δp = 5 psia/ 140 micron	CS	20

	54.	F-05 Therminol Cooler Blower Filter	To filter the solids from air			20
	SPEC	IALIZED EQUIPMENTS				
	35.	R-01 Fluidized Bed Reactor (two)	To reduce SiGi ₄ to Si by Zn	1330.2 Btu/hr, 5.5" dia.	Graphite Lined /SS	20
	56.	FN-01 Furnace (two)	To preheat SiCl ₄	272,966 Btu/hr		20
	57.	B-01 Seed Addition Hopper (two)	To feed Si seed to the reactor		310 SS	20
	58.	B-02 Si Product Hopper (four)	To hold Si product	6 gal	310 SS	20
<u>ာ</u> အ	59.	B-03 %inc Hopper	To hold make up Zinc	40 gal	CS	20
	65.	C-Cl Therminol Cooler Blower	Therminol system air cooler blower	500 acfm fan/electric, 1 1/2 hp., 12-1/2" wheel	rs es	30
	61.	C-02 Scrubber Vent Blower	Suck SiCl ₄ gas for A-01 & A-02	10,000 acfm/electric, 50 hp. 31-1/2" wheel	FRP	1
	62.	E-01 Eductor (two)	SiCl ₄ scrubbing (Scrubber D-05)	20 gpm, $\Delta p = 47.4 \text{ psia/}$ Hydraulic ejector, 1-1/2" NPT	P.V.C.	ī
	63.	EC-01 Electrolysis Cell (six)	To recover Zr. from ZnCl ₂	5,000 ~6,000 $ar_{\tilde{x}}$ cells	Graphite/SS	20
	64.	PW-01 Power Supply	To supply power to electrolysis cell	545,933 Btu/hr.		20
	65.	VP-01 Zinc Vaporizer (two)	To provide zinc vapor to reactor	104,128.8 Btu/hr 13.5" dia. x 32"	Quartz	20

NOTE:

- 1. For the 1000 MT/yr plant, items 3, 4, 33, 46, 47, 48, 49, 61, and 62 are used for waste treatment of distillation wastes (light, heavy) and vent gases.
- 2. In the 50 MT/yr facility, these items are used for hypochlorite manfacture which is not present in the 1000 MT/yr plant.
- 3. For H-11, the operation conditions were changed from 171°F 32°F to 662°F 20°F.
- 4. For H-12, the operations conditions have been chnaged from ΔT = 855°F to 270°F.

TABLE 3.4-7

PRODUCTION LABOR REQUIREMENTS FOR BCL PROCESS - Case A

			Labor	= .
	Section		man-hr/KG Si	(oper/shift)
				
1.	Purification	(1)	0.01402	(2)
2.	Deposition	(II)	0.01402	(2)
3.	Electrolysis	(III)	0.02103	(3)
4.	Waste Treatment	(IV)	0.00701	(1)
5.	Product Handling	(v)	0,00701	(1)
		TOTAL	0.06309	(9)

Note

Manpower estimate for production labor requirements based on:

- Dividing plant into sections

 type of unit operation
 mark off working area
- 2. Specify work duties required in each section
- 3. Estimate operators required to perform work duties in each section
 - -type of unit operation
 - -size of working area
 - -degree of automation (batch, semi-continuous, continuous, etc.)

3.5 BCL Process for Silicon - Case B (Battelle Columbus Laboratories)

The chemical engineering analysis activity involves a preliminary process design of a plant to produce silicon via the technology under consideration.

The process flowsheet for the Case B of the BCL process to manufacture silicon is shown in Figure 3.5-1. This process consists of several major processing operations of distillation, vaporization, strippping, condensation and a deposition reaction to produce silicon as well as electrolysis to receiver the zinc.

Silicon tetrachloride (SiCl₄), which is the major raw material, is fed to the distillation section for purification, to remove impurities (such as boron and phosphorous). In the deposition section, purified silicon tetrachloride is vaporized and preheated to the reaction temperature, 927°C, before it is introduced into a silicon desition unit, which is a fluidized bed reactor. Zinc vapor produced by a specially designed induction-heated vaporize is also introduced to the reactor at the same temperature for the reaction. The reaction equation to show the silicon deposition is

$$SiCl_4 + 2Zn \rightarrow Si + 2ZnCl_2$$
 (3.5-1)

Silicon granules produced by the deposition reaction, which descend to the bottom of reactor, are cooled and collected in containers. A small amount of silicon seed is fed to the reactor to control the particle size of the silicon product. Zinc chloride and unreacted zinc are recovered and fed to the electrolysis section, while unreacted silicon tetrachloride is recycled to the distillation section.

In the electrolysis section, zinc chloride is reduced to zinc by low voltage (4-b volts) electrolysis cells. Zinc is recycled to the deposition unit, while chlorine gas is collected as the by-product. The deposition and electrolysis sections are purged with intert gas (such as argon). Waste gases from various sections are collected and treated with hydrate lime solution in the waste treatment section.

A process design was performed to obtain data for a cost analysis of a plant to produce silicon by this new technology. The design was based on a plant to produce 1000 metric tons/yr of silicon via the BCL process. In case 3, the process contains one deposition reactor and two electrolysis cells as compared with two deposition reactors and six electrolysis cells for Case A which was reported earlier.

The detailed status sheet for the process design package is shown in TAble 3.5-1 and is representative of the various sub-items that make up the activity. The summarized results for the preliminary process design are presented in a tabular format to make it easier to locate items of specific interest.

The guide for these tables is given below:

Base Case Conditions-----Table 3.5-2
Reaction Chemistry------Table 3.5-3
Raw Material Requirements-----Table 3.5-4
Utility Requirements-----Table 3.5-5
Major Process Equipment-----Table 3.5-6
Production Labor Requirements-----Table 3.5-7

The process design provides detailed data for raw materials, utilities, major process equipment and production labor requirements which are necessary for polysilicon production.

	Prel. Process Design Activity	Status	Prel. Process Design Activity Status
1.	Specify Base Case Conditions 1. Plant Size 2. Product Specifics 3. Additional Conditions	•	7. Equipment Design Calculations 1. Storage Vessels 2. Unit Operations Equipment 3. Process Data (P, T, rate, etc.) 4. Additional
	Define Reaction Chemistry 1. Reactants, Products 2. Equilibrium Process Flow Diagram	•	8. List of Major Process Equipment 1. Size 2. Type 3. Materials of Construction
268	1. Flow Sequence, Unit Operations 2. Process Conditions (T, P, etc.) 3. Environmental 4 Company Interaction (Technology Exchange)		8a. Major Technical Factors (Potential Problem Areas) 1. Materials Compatibility 2. Process Conditions Limitations 3. Additional
4.	Material Balance Calculations 1. Raw Materials 2. Products 3. By-Products		9. Production Labor Requirements 1. Process Technology 2. Production Volume
5.	Energy Balance Calculations 1. Heating 2. Cooling 3. Additional	•	10. Forward for Economic Analysis
6.	Property Data 1. Physical 2. Thermodynamic 3. Additional	•	0 Plan ⊕ In Progress ● Complete

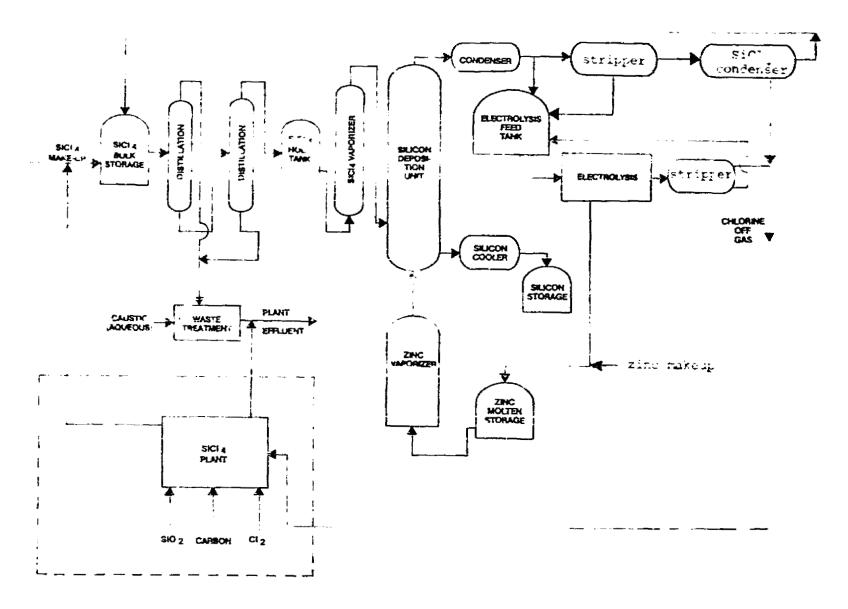


Figure 3.5-1 Process Flow Sheet for BCL Process - Case B

TABLE 3.5-2 BASE CASE CONDITIONS FOR BCL PROCESS (Case B)

1. Plant Size -silicon produced from silicon tetrachloride (TET) -1000 metric tons/yr of silicon -solar cell grade silicon -solid phase product form (granules) 2. Light End Distillation -purification of TET by distillation -remove 4% chlorosilanes as the light end -80°C, 10 psig 3. Heavy End Distillation -purification of TET by distillation -remove 4% impurities as the heavy end -92% over-all yield of TET from both distillations -80°C, 10 psig 4. TET Vaporizer -to supply TET vapor for deposition reactor -by power input (resistance heater) -hold at constant level and constant pressure -164°F 5. Deposition Reactor -reduce TET by zinc to produce silicon -deposit on pure silicon seed -fluid bed -927°C (1700°F, 1 atm) -63% conversion of TET to silicon 6. Reactor Condenser -to condense gases from reactor (ZnCl2, unreacted Zn and SiCl4 gases) -partial condensation -using therminol 66 as the coolant -927°C inlet temperature and 350°C outlet temperature Reactor ZnCl₂ Stripper -work as partial condenser -to condense ${\rm ZnCl}_2{\rm gas}$ from ${\rm SiCl}_4$ gas -operating at the temperature right above ZnCl2 melting point (318°C), 350°C -using therminol 66 as the coolant 8. Call ZnCl, Stripper -operates as partial condenser -to condense ${\rm ZnCl_2}$ gas from ${\rm Cl_2}$ and ${\rm SiCl_4}$ gases -operating at the temperature right above ${\rm ZnCl_2}$ melting point (318°C), 350°C -using therminol 66 as the heat exchange medium

- 9. Reactor SiCl4 Condenser
 - -condense SAC14 gas for recycle
 - -antifreeze as the coolant
 - -350°C inlet temperature, 20°F outlet temperature.
- 10. Electrolysis
 - -electrolytic recovery of Zn from ZnCl2
 - -Cl₂ gas is by product -95% Zn recovery

 - -500°C, approx. 1 atm
- 11. Zinc Vaporizer
 - -to vaporize Zinc
 - -by induction heating
 - -927°C, approx. 1 atm.
- 12. Wastes Treatment
 - -to scrub and neutralize SiCl4 and chlorosilane gases
 - -caustic solution used to neutralize
- 13. Operating Ratio
 - -approximately 80% utilization (on stream time)
 - -approximately 7,000 hr/yr production
- 14. Storage Considerations
 - -feed material (two week supply)
 - -product (two shifts storage)
 - -process (several hours)

TABLE 3.5-3 REACTION CHEMISTRY FOR BCL PROCESS (Case B)

1. Silicon Deposition

$$2Zn + siCl_4 + si + 2ZnCl_2$$

2. Electrolysis

$$ZnCl_2$$
 electroly. $Zn(1) + Cl_2(g)$

3. Waste Treatment

$$SiCl_4 + 2H_2O \xrightarrow{aq} SiO_2 + 4HCl(aq)$$
 $NaOH(aq) + HCl(aq) \rightarrow NaCl(aq) + H_2O$

or

 $Ca(OH)_2(aq) + 2HCl(aq) \rightarrow CaCl_2(aq) + 2H_2O$

3a. Waste Treatment (50 MT/yr anit)

$$Cl_2(g) + 2NaOH(aq) \rightarrow NaOCl(aq) + NaCl(aq) + H_2O$$

Table 3.5-4

Raw Material Requirements for BCL Process (Case B)

	Raw Material	Requirements lb/KG of Silicon
1.	Silicon Tetrachloride, SiCl4	15.33
2,	Zinc, Zn	0.54
3.	Caustic (50%), NaOH(aq)	3.75
or		
	Lime (99%), Ca(OH) ₂	1.75
4.	Argon	3.1 scr*
5.	Nitrogen	7.6 scr*
6.	Chlurine, Cl ₂ (by-product)	11.12

^{*}Estimate from BCL

TABLE 3.5-5 UTILITY REQUIREMENTS FOR BCL PROCESS (Case B)

Requirements/Kg of Silicon Product Utility/Function 25,40 Kw-hr 1, Electricity 1. Low Voltage D.C. for Electro-(20.51)lysis 2. Zinc Vaporizer Induction Wested (4.37)3. Preheat Section of Deposition (1.12) Unit Induction Heated 4. Electrolysis Feed Tank Heater (0.24)(0.10)5. Molten Zinc Storage Heater (0.53)6. SiCl₄ Vaporizer 7. Pumps, Blowers (-2.53)9.67 pounds 2. Steam (50 PSIA) 1. #1 Purification Column Calandria (4.59) 2. #2 Purification Column Calandria (4.30) 3. Caustic Storage Heating (0.29)4. #1 Purificacion Column Preheater (0.49) 3. Cooling Water 34.49 Gallons 1. #1 Purification Column Condenser (16.94) #2 Purification Column Condenser (15.88) 3. Purified Tet Cooler (1.67)10.48 Gallons 4. Process Water 1. Diluent for Waste Treatment (10.48)2.33 MBtu 5. Refrigeration Reactor SiCl₄ Condenser (H-11) SiCl₄ Vent Condenser (H-07) (1.28)(1.10)

TABLE 3.5-6

LIST OF MAJOR PROCESS EQUIPMENT FOR BCL (Case B)

	Equipment	Function	Size/Type	Material of Construction	Capacity Ratio to 1000 MT/yr
PRO	CESS TOWER AND INTERN	IALS			
1.	D-01 Light End Distillation Column	To purify SiCl ₄	8" dia. x 21', packed 13.5'	Column, CS/packing, SS	20
2.	D-02 Heavy End Distillation Column	To purify SiCl ₄	8" dia. x 21', packed 13.5'	Column, CS/packing, SS	20
3.	A-01 Primary SiCl ₄ Vent Scrubber	To scrub SiCl ₄ vent gas	3' día. x 4'4" T/T, 225 gal/flat bottom	FRP	1
4.	A-02 Final SiCl ₄ Vent Scrubber	To scrub SiCl ₄ vent gas	7'6" dia. x 17'4" T/T/ 4 pp trays, Teflon dimister	PRP	1
HEA	AT EXCHANGER				
5.	H-01 L.E. Column Feed Heater	To preheat feed to D-01	2' dia. x 5', 15,013 Btu/hr / external heater	cs	20
6.	H-02 L.E. Column Reboiler	Peboiler of D-01	2' dia. x 3', 51,522 Btu/hr / external heater	cs	20
7.	H-03 L.E. Column Condenser	Total condenser of D-01	47,430 Btu/hr/shell-tube H.E.	cs	20
8.	H-04 H.E. 'olumn Feed Heater	To preheat feed to D-02	2' dia. x 5', 14,331 Btu/hr/external heater	cs	20

9.	H-05 H.E. Column Reboiler	Reboiler of D-02	2' dia. x 3', 56,641 Btu/hr/external heater	CS	20
10.	H-06 H.E. Column Condenser	Total condenser of D-02	52,292 Btu/hr/shell- tube H.E.	CS	20
11.	H-07 SiCl ₄ Vent Condenser	Condense SiCl ₄ from vent gas	38 ft ² , 18,000 Btu/ hr/snell-tube H.E.	CS	20
12.	H-08 SiCl ₄ Vaporizer	To provide SiCl ₄ vapor to reactor	2.75' dia. x 3' T/T, 13,648 Btu/hr/ resistance heater	cs	40
13.	H-09 Reactor Condenser	To condense by products from reactor	14" dia. x 6.4', 126,237.2 Btu/hr	Graphite W/SS shell	40
14.	H-10 Reactor ZnCl ₂ Stripper	To condense ZnCl ₂ gas	12 ft ² , 2,652 Btu/hr/ shell-tube H.E., finned U-tube	316 SS	20
15.	H 11 SiCl4 Condenser	To condense SiCl ₄ gas for recycle	6,401 Btu/hr (x 4.62 = 29.573 Btu/hr)	316 SS	20
16.	H-12 Cell ZnCl ₂ Stripper	To condense ZnCl ₂ vapor	9,841.4 Btu/hr/shell- tube, H.E. (x 0.32)	Inconel 600	20
17.	H-13 Therminol Cooler (cold circuit)	To cool Therminol 66	68 ft ² , 11,000 Btu/hr/ shell-tube H.E., 500 psia	CS	20
18.	H-14 Thermino'. Cooler (hot circuit)	To cool Therminol 66	262 ft ² , 120,000 Btu/hr/ shell-tube H.E., 500 psia		20
19.	H-15 Start-up Heater	Therminol start- up heater	98,950 Btu/hr/U-tube 15', resistance heater	cs	20
20.	H-16 Silicon Product Cooler (two)	To cool the Si product from reactor	5,735 Btu/hr	sic	40

20a.	H-17 Chlorination Cocler		20,000 Btu/hr, Area 200 ft ²	SS	1
20b.	H-18 Cell Gas Cocler		1.08 x 10 ⁵ Btu/hr, Area 1805 ft ²	cs	1
PROC	ESS AND STORAGE VESSE	LS			
21.	T-01 SiCl ₄ Storage Tank	Storage/feed to purification	7' dia. x 16' T/T/ 4,600 gal	cs	20
22.	T-0? SiCl ₄ Emergency Storage Tank	Storage/feed to purification	7' dia. x 16' T/T/ 4,600 gal	CS	20
23.	T-03 L.E. Column Reflux Drum	To hold distillate for reflux	12" dia. x 4'/23 gal	CS	20
24.	T-04 Surge Tank	Surge Tank for D-01 bottom	3' dia. x 4'/200 gal	CS	20
25.	T-05 Sump Tank	Sump for purification unit	3' dia. x 4'/200 gal	CS	20
26.	T-06 H.E. Column Reflux Drum	To hold distillate for reflux	12" dia. x 4'/23 gal	CS	20
27.	T-07 Pure SiCl ₄ Storage Tank	Storage/feed to SiCl ₄ Vaporizer	6' dia. x 10' T/T/ 1900 gal	CS	20
28.	T-08 Electrolysis Feed Tank	Storage/feed ZnCl ₂ to electrolysis cel	50" x 158" x 38"H/ l 7" graphite TH	Graphite/304 SS	26
29.	T-09 Molten Zinc Storage Tank	Storage/feed to Zinc vaporizer	W/heater 68,242 Btu/hr	Graphite/304 SS	20
30.	T-10 Thermino: Nead	Storage Therminol	1.5' dia. x 3.75' T/T/	cs	20

31.	T-11 Therminol Drain Down Tank	To store drained Therminol	2.75' dia. x 3' T/T/ 133 gal	CS	20
32.	T-12 Chlorine Sup- ply Tank	To supply chlorine gas	1 1/2' dia. x 3'/ 37.62 gal	CS	20
33.	T-13 Lime Storage Tank	Storage Lime	12' dia. x 14'6" T/T/ 12,000 gal	FRP	1
PUMP	S WITH DRIVERS				
34.	P-01 Purification Feed Pump	To feed SiCl ₄ to storage tank	30 gpm, 31' head/ centrifugal, 1 1/2 hp	cs	20
35.	P-02 L.E. Column Feed Pump	To supply SiCl ₄ to preheater	28.9 gph, $\Delta p = 72$ psia/ 0.5 hp.	CS	20
36.	P-03 L.E. Column Relux Pump	D-01 Reflux	51.7 gph, Δp = 23 psia/ 0.5 hp.	cs	20
37.	P-04 Surge Tank Pump	To supply SiCl ₄ to H.E. Column	29.4 qph, $\Delta p = 53$ psia/ 0.5 hp.	CS	20
38.	P-05 Sump Pump	To pump Sicl ₄ to emergency tank	30 gpm, 31' head/ centrifugal, 1 1/2 hp.	CS	20
39.	P-06 L.E. Column Bottom Pump	To pump SiCl ₄ to surge tank	29.4 gph, $\Delta p = 53 \text{ psia/} 0.5 \text{ hp.}$	CS	20
40.	P-07 H.E. Column Relux Pump	D-02 Reflux	57.1 gph, $\Delta p = 25 \text{ psia/} 0.5 \text{ hp.}$	cs	20
41.	P-08 H.E. Column Bottom Fire	To pump bottom solution to waste treatment	1.3 gph, $\Delta p = 25$ psia/ 0.5 hp.	cs	20
42.	P-09 SiCl ₄ Vaporizer Feed Pump	To feed SiCl ₄ to Vaporizer	15 gph, 31' head/ 1/2 hp	cs	20

43.	P-10 Reactor Con- denser Circulating Pump	To circulate con- densates	2.4 gpm, 30' head/1/2 hp	Graphite	40
44.	P-11 Cold Circuit Pump (two)	Cold Therminol circulation	20 gpm, 85' head/centri- fugal, 2 hp.	CS	20
45.	P-12 Hot Circuit Pump	Hot Therminol circulation	62 gpm, 85' head/centri-fugal, 4 hp.	cs	20
46.	P-13 Primary Scrubber Recircula- tion Pump	Recirculation for Scrubber A-01	20 gpm, 125' head/ centrigugal, 2.5 hp.	Duriron	1
47.	P-14 Primary Scrubber Lower-loop Recirculating Pump	Circulate solution for Lower-loop of Scrubber A-02	100 gpm, 103' head/ centrifugal, 7 1/2 hp.	Duriron	1
48.	P-15 Primary Scrubber Upper-loop Recirculating Pump	Circulate solution for upper-loop of Scrubber A-02	100 grm, 13' head/ centrifugal, > hp.	Duriron	1
49.	P-16 Make up Lime Metering Pump	Lime make up	0.9 gpm, 25' head/ centrifugal, 1/2 hp.	CS	1
FILT	ERS				
50.	F-01 L.E. Column Feed Filter	Remove solids	29 gph, Δp = 5 psia/ 140 micron	cs	20
51.	F-02 L.E. Column Reflux Filter	Remove solids	30 gph, Δp = 5 psia/ 140 micron	cs	20
52.	F-03 H. E. Column Feed Filter	Remove solids	52 gph, Δp = 5 psia/ 140 micron	cs	20
53.	F-04 H.E. Column Relfux Filter	Remove solids	31 gph, Δp = 5 psia/ 140 micron	cs	20

	54.	F-05 Therminol Cooler Blower Filter	To filter the solid from air	s		20
	SPEC	IALIZED EQUIPMENTS				
	55.	R-01 Fluidized Bed Reactor (two)	To reduce SiCl ₄ to Si by Zn	1830.2 Btu/hr, 6.5" dia.	Graphite Lined /SS	40
	56.	FN-01 Furnace (two)	To preheat SiCl ₄	272,966 Btu/hr		40
	57.	B-01 Seed Addition Hopper (two)	To feed Si seed to the reactor		310 SS	40
	58.	B-02 Si Product Hopper (four)	To hold Si product	6 gal	310 SS	20
))	59.	B-03 Zinc Hopper	To hold make up Zinc	40 gal	Cs	20
	60.	C-01 Therminol Cooler Blower	Therminol system air cooler blower	500 acfm fan/electric, 1 1/2 hp., 12-1/2" wheel	Cs	20
	61.	C-02 Scrubber Vent Blower	Suck SiCl ₄ gas for A-Ol & A-O2	10,000 acfm/el-ctric, 50 hp. 31-1/2 wheel	FRP	1
	62.	E-01 Eductor (two)	SiCl ₄ scrubbing (Scrubber D-05)	20 gpm, Δp = 47.4 psia/ Hydraulic ejector, 1-1/2" NPT	P.V.C.	1
	63.	EC-01 Electrolysis Cell (six)	To recover Zn from ZnCl ₂	5,000 ~6,000 amp cells	Graphite/SS	60
	64.	PW-31 Power Supply	To supply power to electrolysis cell	545,933 Btu/hr.		20
	65.	VP-01 Zinc Vaporizer (two)	To provide zinc vapor to reactor	104,128.8 Btu/hr 13.5" dia. x 32"	Quartz	40

NOTE:

- 1. For the 1000 MT/yr plant, items 3, 4, 33, 46, 47, 48, 49, 61, and 62 are used for waste treatment of distillation wastes (light, heavy) and vent gases.
- 2. In the 50 MT/yr facility, these items are used for hypochlorite maufacture which is not present in the 1000 MT/yr plant.
- 3. For F-11, the operation conditions were changed from 171°F 32°F to 662°F 20°F.
- 4 For \$12, the operations conditions have been changed from "T = 855°F to 270°F.

TABLE 3.5-7

PRODUCTION LABOR REQUIREMENTS FOR BCL PROCESS (Case B)

			Labor		
	Section		man-hr/KG Si	(oper/shift)	
1.	Purification	(I)	0.01402	(2)	
2.	Deposition	(II)	0.01402	(2)	
3.	Electrolysis	(III)	0,02103	(3)	
4.	Waste Treatment	(IV)	0.00701	(1)	
5.	Product Handling	(V)	0.00701	(1)	
		TOTAL	0 06309	(9)	

Note

Manpower estimate for production labor requirements based on:

- 1. Dividing plant into sections
 - -type of unit operation
 - -mark off working area
- 2. Specify work duties required in each section
- 3. Estimate operators required to perform work duties in each section
 - -type of unit operation
 - -size of working area
 - -degree of automation (batch, semi-continuous, continuous, etc.)

3.6 DCS Process (Dichlorosilane)

The chemical engineering analysis activity involves a preliminary process design of a paint to produce dichlorosilane as a silicon source material by using the technology under consideration.

The process flowsheet for the DCS process to produce dichlorosilane consisting several major processing operations of hydrochlorination, condensation, stripping, distillation and redistribution reaction, is shown in Figure 3.6-1.

Metallurgical grade silicon (M.G.Si) is hydrochlorinated at the presence of hydrogen (H₂) and silicon tetrachloride (SiCl₄) in a fluidized bed reactor. The product stream from the hydrochlorination is cooled. A settler is then used to remove metal impurities. The chlorosilanes - dichlorosilane (DCS), trichlorosilane (TCS) and silicon tetrachloride (TFT) are separated by several distillation units. After separation, the silicon tetrachloride is recycled.

Intermediate in the several dist itation units, the TCS is redistributed to DCS and TET by passing through a fixed bed of catalyst. After redistribution that the matter fed to appropriate distillation unit for separation, and preficution. The reaction equations to produce DCS are shown

$$38iC1_4 + 8i + 21I_2 + 48ie^{-i}$$
 (3.6-1)

$$2SiHC1_3 \stackrel{?}{+} SiH_2C1_2 + SiC$$
 (3.6-2)

A process design was performed to obtain data for a cost analysis of a plant to produce DC by this new technology. The design was based on a paint to produce 2,780 metric tons/yr of DCS which is sufficient to produce 1000 metric tons/yr of silicon without recycle.

The detailed status sheet for the process design package is shown in Table 3.6-1 and is representative of the various subtems that make up the activity. The process design are presented in a tabular format to make it easier to locate stems of emeritic interest. The guide for these tables is given below:

	Base Case Conditions.		Tehle	3.6-2
	Reaction Chemistry		Table	3.6 - 3
	Raw Material Requirem nos.		Pable	3.6.4
٠	Utility Requirements		Table	3,6-5
•	Major Process Equipment		Table	3.6-6
•	Production Labor Requiremen	t 4, j	Table	3.6-7

The process design provides detailed data for raw materials, utilities, major process equipment and production labor requirements which are necessary for polysilicon production.

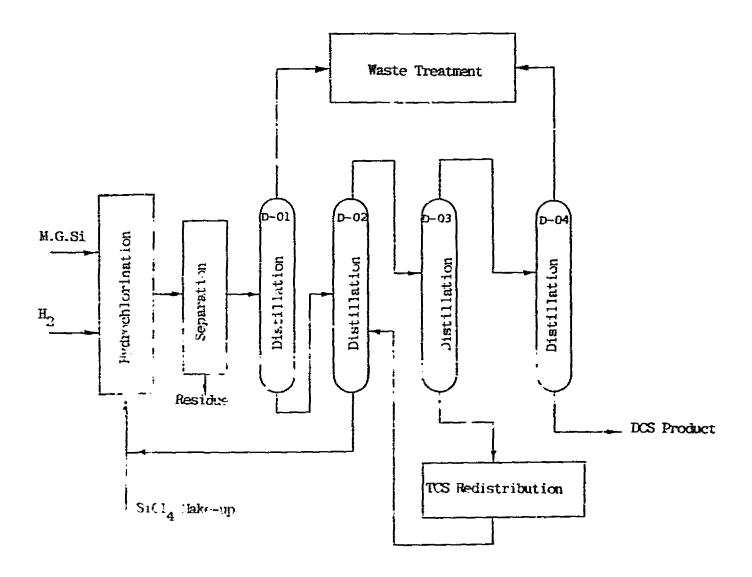


Figure 3.6-1 Process Flowsheet for DCS Process

TABLE 3.6-1 CHEMICAL ENGINEERING ANALYSES: PRELIMINARY PROCESS DESIGN ACTIVITIES FOR DCS PROCESS

	Prel. Process Design Activity	Status	Prel. Process Design Activity Status
	Specify Base Case Conditions 1. Plant Size 2. Product Specifics 3. Additional Conditions	•	7. Equipment Design Calculations 1. Storage Vessels 2. Unit Operations Equipment 3. Process Data (P, T, rate, etc.) 4. Additional
2.	Define Reaction Chemistry 1. Reactants, Products 2. Equilibrium	•	8. List of Major Process Equipment 1. Size 2. Type
3.	Process Flow Diagram 1. Flow Sequence, Unit Operations 2. Process Conditions (T, P, etc.) 3. Environmental 4. Company Interaction (Technology Exchange)	•	3. Materials of Construction 8a. Major Technical Factors (Potential Problem Areas) 1. Materials Compatibility 2. Process Conditions Limitations
4.	Material Balance Calculations 1. Raw Materials 2. Products 3. By-Products	•	9. Production Labor Requirements 1. Process Technology 2. Production Volume
5.	Energy Balance Calculations 1. Heating 2. Cooling 3. Additional	•	10. Forward for Economic Analysis
6.	Property Data 1. Physical 2. Thermodynamic 3. Additional	•	0 Plan 0 In Progress € Complete

TABLE 3.6-2

BASE CASE CONDITIONS FOR DCS PROCESS

- 1. Plant Size
 - -Dichlorosilane produced from m.g. silicon and silicon tetrachloride.
 - -2,895 lb/hr dichlorosilane (enough to support 1000 metric tons/year of silicon production)
 - -9,780 metric tons/yr of DCS capacity
- livdrochlorination
 - Metallurgical grade silicon, hydrogen, and recycle sili-con tetrachloride (TET) used to produce trichlorosilane (TCS)
 - -Copper catalyzed
 - -Fluidized bed

 - -500°C, 514.7 psia -H₂/Cl₂ ration about 2.8 -30% conversion of SiCl₄ to SIHCl₅
- TCS Redistribution Reaction
 - -TCS is redistributed to DCS and TET through catalytic reaction
 - -Catalytic redistribution of TCS with amine function ion exchange resin
 - -Liquid phase 85 psia, 140°F
 - -Conversion from pure TCS feed is about 11% to DCS
- Recycles
 - -Unreacted chlorosilanes and hydrogen are separated by distillation and recycled
- Dichlorosilane Purification
 - -Final purification by distillation
 - -Designed to remove trace impurities (BpHg, example)
- Operating Ration
 - -Approximately 85% utilization (on stream time) -Approximately 7446 hour/year production
- Storage Consideration
 - -Feed materials (several week supply, approx. 1 month)
 - -Product (two shifts storage)
 - -Process (several hours to 1 shift)

TABLE 3.6-3

REACTION CHEMISTRY FOR DCS PROCESS

1. Hydrochlorination Reaction

2. Redistribution Reaction

3. Waste Treatment (representative - overall)

$$SiH_2Cl_2 + Ca(OH)_2 \xrightarrow{aq} SiO_2 + CaCl_2 + 2H_2O$$

 $SiHCl_3 + 1.5Ca(OH)_2 \xrightarrow{aq} SiO_2 + 1.5CaCl_2 + 2H_2O$
 $SiCl_4 + 2Ca(OH)_2 \xrightarrow{aq} SiO_2 + 2CaCl_2 + 2H_2O$

Note:

- 1. Reaction 1 product contains $\rm H_2$, $\rm HCl$, $\rm SiCl_4$, $\rm SiHCl_3$, $\rm SiH_2Cl_2$ (trace), other trace chlorides
- 2. Reaction 2 product contains SiCl_3 , SiCl_4 , $\mathrm{SiH}_2\mathrm{Cl}_2$, $\mathrm{SiH}_3\mathrm{Cl}$

TABLE 3.6-4
RAW MATERIAL REQUIREMENTS FOR DCS PROCESS

	Raw Material	Requirements lb/hr	lb/kg of DCS
1.	M.G. Silicon (Si)	456.6	0.348
2.	Silicon Tetrachloride (SiCl ₄ , make-up)	2609.5	1.987
3.	Liquid Hydrogen	62.5	0.048
4.	Copper Catalyst (Cu)	6.8	0.005
5 .	Hydrate Lime (Ca(OH)2)	310.0	0.236

TABLE 3.6-5
UTILITY REQUIREMENTS FOR DCS PROCESS

	<u>Ut 1</u>	lities/Function	Total Requirements	Requirements per Kg of DCS
1.	Ele	octricity		
	1)	Gas Compressions (96bhp)	217 KW	.165 KW-Hr
	2)	Pumping Liquids (98.5 bhp)	223 KW	.170 KW-Hr
	3)	Filter Drive (1 bhp)	2,3 KW	<u>.002 KW-Hr</u>
			443 KW	,337 KW-Hr
2.	Ste	am (50 psia, saturated)		
	1)	Column Reboiler (12.0MM Btu/hr)	3,850 lb/hr	2.93 lb
	2)	Vaporizer (3.56MM Btu/hr)	12,970 lb/hr	9.88 lb
			16,820 lb/hr	12.81 lb
3.	Coc	oling Water		
	1)	Coolers and Condensers (21.66MM Btu/hr)	1237 gpm	56.51 gal
4.	Pro	cess Water		
	1)	Waste Treatment	6.59 gpm	.301 gal
5,	Fue	01 Oil		
	1)	Direct-Fired Heater (4.71MM Btu/hr)	33.4 gal/hr	.026 gal
	2)	Incineration (1.5 x 10 ⁶ MM Btu/hr)	10.6 gal/hr	.008 gal
			44.0 gal/hr	.034 gal

TABLE 3.6-6

LIST OF MAJOR PROCESS EQUIPMENT FOR DCS PROCESS

		Equipment	Function	Duty/Type	Size	Material of Construction	
		Reactors				<u> </u>	
291	1.	R-01 Hydrochlo- rination Reactor	Hydrochlorination of m.g. Si & SiCl ₄	32,200lb/hr Feed/Fluid. bed	8.54' diam. x 9.75', and 12.25', 30° cone	316SS	
	2.	R-02 TCS Redis- tribution Reactor	Conversion of TCS to DCS	31,000lb/hr Feed/Fined Bed, catalyst	2' diam. x 36' 320 psia	31688	
	3.	R-03 Waste Neutra- lizer	Waste Treatment	agitated vessel	3' diam. x 20' 14.7 psia	31688	
	4.	R-04 Waste Com- buster	To incinerate waste vapors	25 SCFM Vapor/Combustion	3' x 3' x 9' 14.7 psia	CS/Brick	
		Distillation Columns					
	5.	D-01 Crude TCS Stripping Column	To remove inert gases	31,217lb/hr of feed	24" diam. x 20' tall with 10 sieve plates	cs	
	6.	D-02 TCS/STC Dis- tillation Column	To remove STC at bottom	62,208.41b/hr of feed	5.2' diam. x 68'tal with 29 sieve plate:		
	7.	D-03 DCS/TCS Dis- tillation Column	To remove DCS at distillates	34,001.21b/hr of feed	4.9' diam. x 102'ta with 46 sieve plates		
	8.	D-04 DCS Distil- lation Column	To purify DCS	3,009.81b/hr of feed	1.1' diam. x 70' ta with 40 sieve plates	02000	

TABLE 3.6-6 (continued)

TABLE 3.6-6 (Continued)						
		Equipment	Function	Duty/Type	Size	Material of Construction
		Tanks & Bins				
292	9.	B-01 Silicon Sto- rage Bin with Feed Lock	To store and feed m.g. Si to reactor	1 week storage/Ver- tical, with Feed Lock	7'diam. x 22', 60°cone	cs
	10.	T-01 Residue Set- ting Tank	To separate unre- acted solid resi- dues	Vertical	8' diam. x 16', 515 psia	31688
	11.	T-02 Residue With- draw Tank	To remove unreacted solid residues	Vertical	3' diam. x 6', 515 psia	31688
	12.	T-03 Hydrogen Se- paration Tank	To separate H ₂ gas from chlorosilanes	Vertical, mesh pad	3.75' diam. x 11.25 515 psia	', cs
	13.	T-04 Crude TCS Storage Tank	To store crude TCS	8 hr. storage, Horizontal	12'diam. x 33', 100 psia	CS
ស៊	14.	T-05 TCS Stripper Reflux Drum	Reflux drum for D-01 column	30 min. storage, Vertical	2' diam. x 3.5', 90 psia	CS
	15.	T-06 TCS/STC Dis- tillation Ref vx Drum	Reflux drum for D-02 column	10 min. storage, Vertical	4.5' diam. x 19', 55 psia	CS
	16.	T-07 STC Storage Tank	To store STC	6 hr. storage, Rorizontal	10.25' diam. x 3.05 15 psia	', CS
	17.	T-08 DCS/TCS Dis- tillation Reflux Drum	Reflux drum for D-03 column	10 min. storage, Vertical	4' diam. x 13', 320 psia	CS
	18.	T-09 DCS Distil- lation Reflux Drum	Reflux drum for D-04 column	10 min. storage, Vertical	1.5' diam. x 4.25', 125 psia	31688
	19.	T-10 DCS Storage Tank	To store purified DCS	8 hr. storage, Horizontal	6' diam. x 16.75', 125 psia	31688

TABLE 3.6-6 (continued)

		TABLE	3.6-6 (continued)		36-4-mi3
	Equipment	Function	Duty/Type	Size	Material Construction
20.	T-11 Flue Gas Se- paration Tank	To separate flue gas from lime solu- tion	Vertical tank with mesh	2' diam. x 5'	CS
21.	T-12 Lime Solu- tion Preparation Tank	To prepare lime solution	8 hr. storage, Ver- tical, open top	5' diam. x 9.5'	cs
22.	T-13 Waste Fil- trate Storage Tank	To store waste filtrate	4 hr. storage, vertical	5' diam. x 8'	cs
	Heaters & Heat Exch	angers			
23.	H-01 Crude TCS Condenser	To condense chlo- rosilanes	8.4MM Btu/Hr. Shell- Tube H.E.	1211 ft ² , 515 psia	31688
2 24. 3	H-02 H ₂ Gas Pre- heater ²	To preheat H ₂ Gas for chiorination	500°C discharge, Direct-fired heater	2.59MM Btu/hr, 515 psia	CS/316SS
25.	H-03 STC Vaporizer	To vaporize and superheat STC for chlorination	3.56MM Btu/hr, Kettle	573 ft ² , 515 psia	316SS
26.	H-04 Stripper Condenser	Partial condenser for D-01 column	86,700 Btm/hr Shell-tube H.E.	30 ft ² , 90 psia	cs
27.	H-05 Stripper Reboiler	Stripper reboiler of D-01 column	0.91MM Btu/hr, Kettle	40 ft ² , 95 psia	CS
28.	H-06 TCS Condenser	To condense TCS vapor of D-02 column	7.9MM Btu/hr, Shell-tube H.E.	2,358 ft ² , 55 psia	31655
29.	H-07 TCS/STC Re- boiler	Reboiler for D-02 TCS/STC Distilla- tion column	8.25MM Btu/hr, Kettle	318 ft ² , 55 psi a	CS
30.	H-08 STC Heat Exchanger	STC Cooling and Heating	0.824MM Btu/ar, Liq- liq. heat exchanger	742 ft ² , 55 psia	316SS/CS

	TABLE 3.6-6 (continued)				34 2 2 3 3 3 5 7	
		Equipment	Function	Duty/Type	Size	Material of Construction
	31.	H-09 DCS Condenser	To condense DCS Vapor from D-03 Column	3.2MM Btu/hr Shell- Tube H.E.	328 ft ² , 320 psia	31688/CS
	32.	H-10 DCS/TCS Reboiler	Reboiler for DCS/ TCS distillation column, D-03	2.7MM Btu/hr, Kettle	252 ft ² , 320 psia	CS
	33.	H-11 TCS Cooler	To cool TCS before redistribution reaction	1.32MM Btu/hr Shell-Tube H.E.	78.6 ft ² , 85 psia	31688
	34.	H-12 DCS Distil- lation overhead condenser	To condense over- head of D-04 column	0.183MM Btu/hr Shell-Tube H.E.	23.4 ft ² , 355 psia	3168S/CS
S)	35.	H-13 DCS Distil- lation Reboiler	Reboiler of DCS Dis- tillation column,D-04		6.8 ft ² , 355 psia	31688/C8
94	36.	H-14 Waste stream cooler	To cool waste stream in waste treatment	0.5MM Btu/hr Shell-Tube H.E.	125 ft ² , 60 psia	316SS/CS
	37.	H-15 STC Super- heater	To heat STC bef re hydrochlorination	500°C discharge temp. Direct-fired heater	2.12MM Btu/hr, 515 p	osia 316SS/CS
	38.	H-16 H ₂ Compressor Intercooler	To cool H ₂ gas bet- ween compression stages	70,000 Btu/hr Shell-Tube H.E.	67.7 ft ² , 90 psia	316SS/CS
		Compressors and Pum	<u>os</u>			
	39.	C-01A Hydrogen Feed Compressor. First stage	Compression of recycle and make-up	187 SCFM/Recip.comp.	38bhp. discharge press. 87 psia	cs
	40.	C-01B Hydrogen Feed compressor, Second stage	Compression of recycle and make-up H ₂ gas	187 SCFM/Recip.comp.	41bhp., discharge press. 515 psia	CS

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TABLE 3.6-6 (continued)

		THOME 5.	6-6 (constitued)		
	Equipment	Function	Duty/Type	Size	Material of Construction
41.	C-02 Hydrogen Circulation Com- pressor	Compression of recycle H ₂ gas	2,833 SCFW/centri-	17bhp., ΔP=30 psi	cs
42.	P-01 Feed Tank Blower	To load silicon to its storage bin	Pnumatic transport/ centrifugal blower	939ACFM, 32bhp	CS
43.	P-02 Settling Tank Circulation Pump	Circulation and to support ejector	100 gpm centrifugal	37' Head, 1.75bhp	31688
44.	P-04 TCS Reflux Pump	Pumping TCS for D-02 reflux and feed to D-03	58 gpm centrifugal	12bhp, discharge press. 320 psia	CS
45.	P-05 STC Feed Pump	Pumping STC to Hydro- chlorination reactor	-42.2 gpm centrifugal	15bhp, discharge press. 500 psia	CS
9 46. 9	P-06 DCS Reflux Pump	Pumping DCS for D-03 reflux and feed to D-04	6.5 gpm centrifugal	łbhp, discharge press. 355 psia	31688
47.	P-08 DCS Puri- fication Discharge Pump	To withdraw impuri- ties from DCS Puri- fication unit	2.6 gpm centrifugal	thp, discharge press. 355 psia	316\$\$
48.	P-09 DCS Pump	To pump pure DCS	290 gpm centrifugal	34 bhp	31688
49.	P-10 Waste Solu- tion Pump	To feed slurry to filter	12.5 gpm centrifugal	1.25 bhp	Cast Iron
50.	P-11 Lime Solu- tion Circulation Pump	To circulate lime solution to neutra-lizer	12.5 gpm centrifugal	1.25 bhp	Cast Iron
51.	P-12 Fresh Lime Solution Pump	To supply frash lime solution	6.5 gpm centrifugal	0.75 bhp	Cast Iron

TABLE 3.6-6 (continued)

	TABLE 3.6-6 (Continued)				
	Equipment	<u>Function</u>	Duty/Type	Size	Material of Construction
	Miscellaneous				
52.	F-01 Silicon Dust Filter	To retain m.g. silicon dust	Gas-Solid/Bag	20 ft ² x 5μ	CS/cloth
53.	F-02 Waste Slurry Filter	To remove waste sludge	12.5 gpm rotary filter	2 ft ²	CS/cloth
54.	S-01 Silicon Feed Cyclone	To feed m.g. sili- con to storage bin	6" w.c.ΔP	940ACFM	31688
55.	E-01 Quench Contact Ejector	To withdraw and cool effluent of hydrochlorination	6" w.c. suction	100 gpm 134 ACFM	316SS
56.	E-02 Flue Gas Ejector	To withdraw flue gas from waste gas com- bustion		100 gpm 1 SCFM	CS

TABLE 3.6-7
PRODUCTION LABOR REQUIREMENTS FOR DCS PROCESS

Labor

	Section	man-hr/KQ DCS	(oper/shift)
1.	Hydrochlorination	0.001294	(2)
2,	Purification/Re- distribution	0.00194	(3)
з.	Waste Treatment	0.000847	(1)
	TOTAL	0.003882	(6)

Note

Manpower estimate for production labor requirements based on

- Dividing plant into sections
 -type of unit operation
 -mark off working area
- 2. Specify work duties required in each section
- 3. Estimate operators required to perform owrk duties in each section
 - -type of unit operation
 - -size of working area
 - -degree of automation (batch, semi-continuous, etc.)

4. ECONOMIC ANALYSES

4.1 SiI_A Decomposition Process

The economic analysis activity involves a cost analysis of the process under consideration for the production of silicon. The cost analysis for the particular technology is based on process design results, such as requirements for raw materials and major process equipment necessary to produce the product, from the chemical engineering analysis activity. Primary results issuing from the economic analysis include plant capital investment and product cost which are useful in identification of those processes showing promise for meeting project cost goals.

The cost analysis results for producing silicon by the Sil, decomposition process are presented in Table 4.1-1 including costs for raw materials, labor utilities and other items composing the product cost (total cost of producing silicon). The tabulation summarizes all of these items to give a total product cost without profit of \$44.64 (1975 dollars) and \$62.50 (1989 dollars) per kg. This product cost without profit includes direct manufacturing cost, indirect manufacturing cost, plant overhead and general expenses.

The product cost represents all cost associated with producing silicon. On top of these costs a producing company will includes some profit. The sales price of the product silicon will actually be the sum of the product cost and a profit for the company. The profit is usually measured in terms of rate of return on the capital investment that the company spent in going into the polysilicon business. Two profitability methods which are commonly used are the return on original investment (per cent ROI) and discounted cash flow rate of return (per cent DCF).

The cost and profitability analysis summary for this process are presented in Table 4.1-2. The sales price of polysilicon at various rates of return for both profitability methods (per cent KOI and DCF) is shown in the lower half of the table. The results indicate a sales price of \$71.48 per kg of silicon (1980 dollars) at 5 per cent DCF return on investment.

These cost and profitability results for the SiI_A decomposition process indicate that this new technology for producing polysilicon does not show promise for meeting the cost goal of \$14 per kg of silicon material (1980 dollars) for solar cells.

The detailed results for the economic analysis are presented in a tabular format to make it easier to locate cost items of specific interest. The guide for the tabular format is given below:

•	Preliminary Economic Analysis Activities Table	4.1-3
•	Process Design Inputs Table	4.1 - 4
	Base Case Conditions	
•	Raw Material Cost	4.1 - 6
•	Utility Cost	4.1 - 7
•	Major Process Equipment Cost	4.1 - 8
•	Production Labor Cost	4.1 - 9
٠	Plant Investment	4.1-10
•	Total Product Cost	4.1 - 11

		Cost \$/Kg of Silicon (1975 dollars)	Cost \$/Kg of Silicon (1980 dollars)
1.	Direct Manufacturing Cost (Direct Costs) Raw Materials Direct Operating Labor Utilities Supervision and Clerical Maintenance and Repairs Operating Supplies Laboratory Charge	23.48	32.67
2.	Indirect Manufactuing Cost (Fixed Cost) Depreciation Local Taxes Insurance	12.17	17.04
3.	Plant Overhead	3.17	4.44
4.	General Expenses	5.82	8.15
5.	Product Cost Without Profit	44.64	62.50

TABLE 4,1-2

COST AND PROTITABILITY	BIRYIANA YO	SUMMARY	FOR	SiI,	DECOMPOSITION	PROCESS
------------------------	-------------	---------	-----	------	---------------	---------

		*4	
1. 2.	Process	\$iI, Decompos	ition Process Tons/year
Э.	Plant Product	· · · Silicon	
4.	Product Form	Silicon Ingot	s (Rods)
5.	Plant Investment	\$107,600,000/ (1975 dollars)	\$150,650,000 (1980 dollars)
	Fixed Capital Working Capital (15') Total	\$ 93.57Mega \$ <u>14.03Mega</u> \$107.60Mega (1975 dollars)	\$131.00 Mega \$ 19.65 Mega \$150.65 Mega (1980 dollars)
		(TA\2 COTTSER)	(1880 dotture)

6. Return on Original Investment, after taxcs (%ROI)

		\$/Kg of Silicon (1975 dollars)	\$/Kg of Silicon (1980 dollars)
0%	ROI	44.64	62.50
5%	ROI	54.60	76.45
10%	ROI	64.57	90.40
15%	ROI	74.53	104.35
20%	ROI	84.49	118.30
	ROI		132.25
	ROI	104.42	146.19
40%	ROI	124.35	174.09

7. Discounted Cash Flow Rate of Return, after taxes (* DCF)

		\$/Kg of Silicon (1975 dollars)	\$/Kg of Silicon (1980 dollars)	
0%	DCF	44.64	62.50	
5%	DCF	51.05	71.40	
10%	DCF	58.11	81.36	
15%	DCF	65.74	92.04	
20%	DCF	73.84	103.38	
25%	DCF	82.34	115.28	
30%	DCF.,	91.16	127.63	
40%	DCF	109.50	153.31	

Based on 10 year project life and 10 year straight line depreciation.

TABLE 4.1-3

ECONOMIC ANALYSES: PRELIMINARY ECONOMIC ANALYSIS ACTIVITIES FOR SII4 Decomposition Process

	Prel. Process Economic Activity	Status	Prel. Process Economic Activity	<u>e-atus</u>
1.	Process Design Inputs 1. Raw Material Requirements 2. Utility Requirements 3. Equipment List	•	 6. Production Labor Costs 1. Base Cost Per Man Hour 2. Cost/Kg Silicon Per Area 3. Total Cost/Kg Silicon 	•
	4. Labor Requirements	•	7. Estimation of Plant Investment	•
2.	Specify Base Case Conditions 1. Base Year for Costs 2. Appropriate Indices for Costs 3. Additional	• •	 Battery Limits Direct Costs Other Direct Costs Indirect Costs Contingency Total Plant Investment 	
3.	Raw Material Costs 1. Base Cost/Lb. of Material 2. Material Cost/Kg of Silicon	. •	(Fixed Capital) 8. Estimation of Total Product Cost	•
	3. Total Cost/Kg of Silicon	•	 Direct Manufacturing Cost Indirect Manufacturing Cost 	•
4.	Utility Corts 1. Base Cost for Each Utility 2. Utility Cost/Kg of Silicon 3. Total Cost/Kg of Silicon	• • •	 Plant Overhead By-Product Credit General Expenses Total Cost of Product 	•
5.	Major Process Equipment Costs 1. Individual Equipment Cost 2. Cost Index Adjustment	•	0 Plan 0 In Progress	
			● Complete	

TABLE 4.1-4

PROCESS DESIGN INPUTS FOR SII, Decomposition Process

- 1. Raw Material Requirements
 -Silicon tetrachloride, zinc, lime, argon and nitrogen
 -see table for "Raw Material Cost"
- 2. Utility -electricity, steam, cooling water and process water -see table for "Utility Cost"
- 3. Equipment List
 _41 plus picces of major process equipment
 _process vessels, heat exchangers, reactor, etc.
- 4. Labor Requirements -production labor for purification, deposition, etc. -see table for "Production Labor Cost"

TABLE 4.1-9 BASE CASE CONDITION FOR SII4 Decomposition Process

1. Capital Equipment

- -January 1975 Cost Index for Capital Equipment Cost
- -January 1975 Cost Index Value = 430

2. Utilities

- -Electrical, Steam, Cooling Water, Nitrogen
- "January 1975 Cost Index (U. S. Dept. Labor)
- -Values determined by literature search and summarized in cost standardisation work

3. Raw Material Cost

- -Chamical Marketing Reporter
- -January 1975 Value
- -Raw Material Cost Index for Judentrial Chemicals
- -1975 Cost Index Value = 105 / 1905 lesale Price Index, Producer Price Index)

4. Labor Cost

- -Average for Chemical Petroleus coal and Allied Industries (1975)
- -Skilled \$6.90/hi

5. Update to 1980

- "historically cited 1975 dollar (USA project)
- -DOE decision to change to 1989 dellars (JPL, 6/22/79)
- -reports to reflect both 1974 and 1980 dollars (JPL, 6/22/79)
- -inflation factor of 1.4 to be used (JPL, 6/22/79)

TABLE 4.1-6

RAW MATERIAL COSTS FOR SiI₄ DECOMPOSITION PROCESS

	Raw Material	Requirement lb/Kg of Silicon	\$/1b of Material	Cost \$/Kg of Silicon
1.	Metallurgical Si	2.6194	. 454	1.188
2.	Iodine*	.7495	2,59	1.94
3.	Iodine*	6.746	.20	1.35
			TOTAL CO	ST 4.48

^{*} SiI_4 wastes are recovered as iodine for recycle at \$.20/pound. Assuming 10% losses in this step, 10% of total iodine must be purchased at \$2.59/pound.

TABLE 4.1-7

UTILITY COST FOR SII, DECOMPOSITION PROCESS

Utility	Requirements in Kw-hr/ Kg of Silicon	Cost of Utility	Cost \$/Kg of Silicon
Electricity*	218.62	\$.0324/Kw-hr	7.08

^{*} For costing purposes only. Actual utilities would involve cooling water, steam, etc.

	Equipment	Purchased Cost, \$M
1.	Purified SiI ₄ Hold Tank	63.80
2.	Liquid Reactor Overheads Storage	60.50
3.	Silicon Product Storage	6.72
4.	Liquid lodine Storage	4.54
5.	Sil, Bulk Storage	67.19
6.	Metallurgical Silicon Storage	8.06
7.	Feed Tank for Purification Column 2	6.89
8.	Sil ₄ Vaporizer	7.87
9.	Silicon Cooler	3.16
10.	Deposition Condenser	525.4
11.	Separation Column Preheater	3.48
12.	Separation Column O/H Condenser	15.45
13.	Separation Column Calandria	12.19
14.	Separation Column O/H After Cooler	3.09
15.	Iodine Vaporizer	7.86
16.	Iodination O/H Condenser	50.22
17.	Separation Column Bottoms After Cooler	3.48
18.	Tet Purification Preheater	7.73
19.	Tet Purification Column 1 O/H Condenser	15.45
20.	Tet Purification Column 1 Calandria	12.39
21.	Tet Purification Column 2 O/N Condensar	15.45
22.	Tet Purification Column 2 Calandria	12.06
23.	Tet Purification After Cooler	7.73

TABLE 4.1-9 (Continued)

	Equipment	Purchased Cost, \$M
24.	Deposition De-superheaters (6 units)	38.68
25.	Purified Sil Pump	1.76
26.	Deposition Compressor System	12090.0
27.	I2/SiI4 Liquid Pump	1.76
28.	I2/SiI4 Separation Column O/H Pump	1.76
29.	12/Sil4 Separation Column Bottoms Pump	2.15
30.	Iodine Pump	2.15
31.	SiI _A Pump	1.76
32.	Tet Purification Column 1 O/H Pump	1.76
33.	Tet Purification Column 1 Bottoms Pump	2.12
34.	Tet Purification Column 2 Feed Pump	1.25
35.	Tet Purification Column 2 O/H Pump	1.76
36.	Tet Purification Column 2 Bottoms Pump	2.09
37.	Deposition Units (6 units)	1922.0
38.	Iodination Reactor	10.75
39.	I ₂ /SiI ₄ Distillation Column	16.50
40.	Tet Purification Column 1	47.25
41.	Tet Purification Column 2	24.44
	TOTAL PURCHASED COST	15090.65

PRODUCTION LABOR COSTS
Sil DECOMPOSITION PROCESS

	Skilled Man-Hrs/Kg Silicon	Semiskilled Man-Hrs/Kg Silioon	\$Kg Si
1	.008395	-	.05793
2	.01314	-	.09067
3	.008395	-	.05793
4	.008395	-	.05793
5	.00584	-	.040296
6	.008395	-	.05793
7	.00584	-	.040296
8.		.01314	.064386
9.	-	.01314	.064386 \$.5319/Kg

Based on labor costs of \$6.90 skilled, \$4.90 semiskilled.

NOTES

TABLE 4.1-10

ESTIMATION OF PLANT INVESTMENT FOR SII4 DECOMPOSITION PROCESS

		Investment \$1000
1.	DIRECT PLANT INVESTMENT COSTS 1. Major Process Equipment Cost 2. Installation of Major Process Equipment	15,090.0 6,488.7
	3. Process Piping, Installed 4. Instrumentation, Installed	11,166.6 2,867.1
	5. Electrical, Installed 6. Process Buildings, Installed	1,509.0 1,509.0
la.	SUBTOTAL FOR DIRECT PLANT INVESTMENT COSTS (PRIMARILY BATTERY LIMIT FACILITIES)	38,630.4
2.		B 445 5
	 Utilities, Installed General Service, Site Development, Fire Protection, etc. 	7,243.2 1,810.8
	 General Buildings, "frees, Shops, etc. Receiving, Shipping Facilities 	2,112.6 3,168.9
2 a .	SUBTOTAL FOR OTHER DIRECT PLANT INVESTMENT COSTS (PRIMARILY OFFSITE FACILITIES OUTSIDE BATTERY LIM	14,335.5 ITS)
3.	TOTAL DIRECT PLANT INVESTMENT COST, la + 2a	52,965.9
4.	INDIRECT PLANT INVESTMENT COSTS	
	1. Engineering, Overhead, etc.	8,299.5
	2. Normal Cont. for Floods, Strikes, etc.	10,713.9
4a.	TOTAL INDIRECT PLANT INVESTMENT COST	19,013.4
5.	TOTAL DIRECT AND INDIRECT PLANT INVESTMENT COST, 3 + 4a	71,979.3
б.	OVERALL CONTINGENCY, % of 5	21,593.8
7.	FIXED CAPITAL INVESTMENT FOR PLANT, 5 + 6	93,573.(1975 dollars) x 1.4 inflation 131,002.3(1980 dollars)
		137'005'3 (1300 dOTTUES)

TABLE 4.1-11
ESTIMATION OF TOTAL PRODUCT COST FOR SII4 DECOMPOSITION PROCESS

		\$/KG of Si	
1.	Direct Manufacturing Cost (Direct Charges)		
	1. Raw Materials	4.48	
	2. Direct Operating Labor	.58	
	3. Utilities	7.08	
	4. Supervision and Clerical	.08	
	5. Maintenance and Repairs	9.36	
	6. Operating Supplies	1.87	
	7. Laboratory Charge	.08	
2.	Indirect Manaufacturing Cost (Fixed Charges)		
	1. Depreciation	9.36	
	2. Local Taxes	1.87	
	3. Insurance	.94	
3.	Plant Overhead	3.17	
4.	By-Product Credit		
4a.	Total Manufacturing Cost, 1 + 2 + 3 + 4	38.82	
5.	General Expenses		
	1. Administration	2.33	
	2. Distribution and Sales	2.33	
	3. Research and Development	1.16	
6.	Total Cost of Product, 4a + 5	44.64 (1975 dolla <u>x 1.4</u> inflation	
		62.50 (1980 dolla	rs)

4.2 Conventional Polysilicon Process (Siemen's Technology)

The economic analysis activity for the conventional polysilicon process involves a cost analysis for the production of silicon via the Siemen's technology. In the Siemen's technology, trichlorosilane (TCS) is used as the feed source material for semiconductor grade silicon.

Since several existing plants producing semiconductor grade polysilicon in the United States were constructed in the 1960's, the cost analysis is based on a poly plant constructed in the 1960's (1965 or earlier). Operating costs for the plant are applicable to the time period of interest (such as 1975 and 1980).

The cost analysis results for producing silicen by the conventional Siemen's process are presented in Table 4.2-1 including costs for raw materials, labor, utilities and other items composing the product cost (total cost of producing silicon). The tabulation summarizes these items to give a total product cost without profit of \$35.5? If This product cost without profit includes dollars) per kg. This product cost without profit includes direct manufacturing cost, indirect manufacturing cost, plant overhead and general expenses. The range for product cost reflects low and high electrical costs (1.5-30/kw hr for 1975 and 2.1-4.20/kw hr for 1980).

Electrical costs vary with location different costs for different states and different costs for different regions in the same state). However, the range (1.5-3c/kw hr) and intermediate value (2.25c/kw hr) for 1975 are considered representative based on a recent plant site survey listing industrial power cost in the USA. With respect to the intermediate value, the survey indicated the following typical electrical cost for industrial power: Michigan (2.48), Arizona (2.27), Missouri (2.05) and Texas (1.49).

In Table 4.2-1, the average product cost without profit is given as \$38.41 (1975 dollars) and \$53.77 (1980 dollars) per kg for the conventional polysiticon process. This average product costs corresponds to intermediate electrical costs (2.25¢/kw hr for 1975 and 3.15¢/kw hr for 1980). These costs results for the conventional polysiticon process indicate that this Siemen's technology using trichlorositane to producing polysiticon does not show promise for meeting the cost goal of \$14 per kg of silicon material (1980 dollars) for solar cells.

When solar cells come into more widespread use, the capacity of existing polysilicon plants will be exceeded necessitating a new poly plant or plants. Higher capital investment costs will be required for new plants to produce polysilicon by the conventional process. The higher capital investment cost for silicon production in new plants will, of course, appear in higher product cost for polysilicon in terms of increased depreciation, taxes, insurance, etc. The profit will also be higher for a reasonable return on investment for the producing company. Thus, the sales price (product cost with profit) for polysilicon from new plants will be considerably higher than the present price for polysilicon of semiconductor grade produced in existing plants.

The detailed results for the economic analysis are presented in a tabular format to make it easier to locate cost items of specific interst. The guide for the tabular format is given below:

	Preliminary Economic Analysis Activities., Table	4.2-2
	Process Design Inputs	
•	Base Case Conditions	4.2 - 4
•	Raw Material Cost	4,2-5
•	Utility Cost	4.2 - 6
٠	Major Process Equipment Cost,	4.2-7
•	Production Labor Cost	4.2 - 8
٠	Plant InvestmentTable	4.2-9
٠	Total Product Cost	4.2 - 10

TABLE 4.2-1
ESTIMATION OF PRODUCT COST FOR CONVENTIONAL POLYSILICON PROCESS

		Cost \$/Kg c? Silicon (1975 dollars)	Cost \$/Kg of Silicon (1980 dollars)
τ,	Direct Manufacturing Cost (Direct Costs) Raw Materials Direct Operating Labor Utilities Supervision and Clerical Maintenance and Repairs Operating Supplies Laboratory Charge	24.94-30.71	24.92-42.99
2.	Indirect Manufactuing Cost (Fixed Cost) Depreciation Local Taxes Insurance	1.38	1,93
3.	Plant Overhead	3.82	5.35
4.	General Expenses	5,38	7.54
5.	Product Cost Without Profit	35.52-41.29	49.73-57.81
6.	Average Product Cost Without Profit	38,41	53.77

Basis: The above results are based on a plant constructed in the 1960's. (1965) or earlier) which is fully depreciated. The range for product cost without profit relfects low and high electrical costs (1.5~3¢/kw. hr for 1975 and 2.1-4.2¢/kw hr for 1980). The average product cost without profit reflects intermediate electrical costs (2.25¢/kw hr for 1975 and 3.15¢/kw for 1980).

TABLE 4.2-2 ECONOMIC ANALYSES: PRELIMINARY ECONOMIC ANALYSIS ACTIVITIES FOR CONVENTIONAL POLYSILICOM PROCESS

	Prel. Process Economic Activity	Status		Prel. Process Economic Activity	S-atus
1.	Process Design Inputs	•	6.	Production Labor Costs	•
	 Raw Material Requirements 	•		 Base Cost Per Man Bour 	•
	Utility Requirements	•		Cost/Kg Silicon Per Area	•
	3. Equipment List	•		3. Total Cost/Kg Silicon	•
	4. Labor Requirements	•			
			7.	Estimation of Plant Investment	•
2.	Specify Base Case Conditions	•		 Battery Limits Direct Costs 	•
	 Base Year for Costs 	•		2. Other Direct Costs	•
	Appropriate Indices for Costs	•		3. Indirect Costs	
	3. Additional	•		4. Contingency	•
				5. Total Plant Investment	
3.	Raw Material Costs	•		(Fixed Capital)	
3	 Base Cost/Ib. of Material 	•			
П	Material Cost/Kg of Silicon	•	8.	Estimation of Total Product Cost	•
	3. Total Cost/Kg of Silicon	•		l. Direct Manufacturing Cost	•
	· -			2. Indirect Manufacturing Cost	•
4.	Utility Costs	•		 Plant Overhead 	•
	l. Base Cost for Each Utility	•		4. By-Product Credit	•
	2. Utility Cost/Kg of Silicon	•		5. General Expenses	•
	Total Cost/Kg of Silicon	•		6. Total Cost of Product	•
5.	Major Process Equipment Costs	•			
	 Individual Equipment Cost 	•			
	Cost Index Adjustment	•			
				0 Plan	
				• In Progress	
				● Complete	

TABLE 4.2-3

PROCESS DESIGN INPUTS FOR · CONVENTIONAL POLYSILICON PROCESS

- 1. Raw Material Requirements
 - -M.G. silicon, anhydrous HCl, caustic, hydrogen, silicon tetrachloride (by-product; -see table for "Raw Material Cost"
- 2. Utility
 - -electrical, steam, cooling water, etc.
 - -see table for "Utility Cost"
- 3. Equipment List
 - -63 pieces of major process equipment
 - -process vessels, heat exchangers, reactor, etc. -see table for "Major Process Equipment Cost"
- 4. Labor Requirements
 - -production labor for deposition, vaporization, product handling, etc.
 - -see table for "Production Labor Cost"

TABLE 4.2-4

BASE CASE CONDITIONS FOR CONVENTIONAL POLYSILICON PROCESS

- 1. Capital Equipment
 - -January 1975 Cost Index for Capital Equipment Cost
 - -January 1975 Cost Index Value = 430
- 2. Utilities
 - -Electrical, Steam, Cooling Water, Nitrogen
 - -January 1975 Cost Index (U.S. Dept. Labor)
 - -Values determined by literature search and summarized in cost standardization work
- 3. Raw Material Cost
 - -Chemical Marketing Reporter
 - -January 1975 Value
 - -Other Sources
- 4. Labor Cost
 - -Average for Chemical Petroleum, Coal and Allied Industries (1975)
 - -Skilled \$6.90/hr
 - -Semiskilled \$4.90/hr
- 5. Update to 1980
 - -historically cited 1975 dollars (LSA project)
 - -DOE decision to change to .980 dollars (JPL, 6/22/79)
 - -reports to reflect both 19/3 and 1980 dollars (JPL, 6/22/79)
 - -inflation factor of 1.4 to be used (JPL, 6/22/79)

TABLE 4.2-5

RAW MATERIAL COST FOR
CONVENTIONAL POLYSILICON PROCESS

Ra	w Material	Requirement lb/Kg of Silicon	\$/1b of Material	Cost \$/Kg of Silicon
1.	M.G. Silicon	6.72 (Kg/Kg)	1.0/Kg	6.72
2.	Anhydrous HCl	57.96	.10	5.79
3.	Hydrogen	.828	.96	.79
4.	Caustic (50% NaOH) 53.29	.0382	2.04
5.	siCl ₄ (By Product) 46.12	.135	-6.23 (oredit)
			TOTAL COST	9.11 (1975 dollars) x 1.4 inflation 12.75 (1980 dollars)

TABLE 4.2-6
UTILITY COST FOR CONVENTIONAL
POLYSILICON PROCESS

	Utility	Requirements/Kg of Silicon	Cost of Utility	SPsiiiign
1.	Electricity	384.6 kw-hr	\$.03/kw-hr	\$ 11.54
2.	Steam	152 Pounds	_ *	-
3.	Cooling Water	984.5 Gallons	\$.08/M Gal.	.08
4.	Process Water	320.9 Gallons	\$.35/M Gal.	.11
5.	Refrigerant (-40°F)	42.1 M BTU	\$10.38/MM BTU	.44
6.	Refrigerant (34°F)	22.3 м вти	\$ 3.75/MM BTU	.35
7.	High Temperature Coolant	582 Founds	\$ 2.7/M Pounds	1.57
8.	Nitrogen	349 SCF	\$.50/M SCF	.17
			TOTAL COST	14.26 (1975 dollars) x 1.4 inflation 19.96 (1980 dollars)

NOTES

^{*} All steam produced by cooling jacket on polysilicon rod reactor.

TABLE 4.2-7

PURCHASED COST OF MAJOR PROCESS EQUIPMENT FOR CONVENTIONAL POLYSILICON PROCESS

	Equipment		Purchased Cost, \$M
1.	(T1)	M.G. Silicon Storage Hopper	24.1
2.	(T2)	Liquid HCl Storage Tank	435.96
3.	(T3)	Crude TCS Hold Tank (3)	178.8
4.	(T4)	Waste Hold Tank	14.9
5.	(T5)	TCS Reactor Off Gas Flash Tank	7.2
6.	(T6)	Hydrogen Storage Tank	152.1
7.	(T7)	Polysilic n Storage Space	10.8
8.	(T8)	Tet Storage Tanks (2)	85.2
9.	(T9)	Tet Feed Tanks (2)	57.8
10.	(T10)	TCS Feed Tanks (3)	42.6
11.	(T11)	TCS Storage Tanks (3)	127.8
12.	(T12)	TET/TCS Feed Tanks (3)	54.
13.	(T13)	Caustic Storage Tank	106.7
14.	(T14)	#1 Distillation Condenser Flash Tank	.85
15.	(T15)	Rod Reactor Off Gas Flash Tank	7.2
16.	(Hl)	HCl Vaporizer	2.5
17.	(H2)	TCS Reactor Off Gas Cooler	7
18.	(H3)	TCS Reactor Off Gas Condenser	46.3
19.	(H4)	#1 Scrubber Vapor Heater	.75
20.	(H5)	#1 Distillation Column Condenser	14.
21.	(HG)	#1 Distillation Column Calandria	9.25
22.	(H7)	#2 Distillation Column Condenser	14.6
23.	(BH)	#2 Distillation Column Calandria	11.92
24.	(H9)	#3 Distillation Column Condenser	9.1
25.	(H10)	#3 Distillation Column Calandria	5.8
26	(H11)	TCS Vaporizer	1.8
27.	(H12)	Rod Reactor Off Gas Cooler	49.4
28.	(H13)	Rod Reactor Off Gas Condenser	97.5
29.	(H14)	#2 Scrubber Vapor Heater	5.8
30.	(H15)	Liquid Recycle Heater	2.3
31.	(H16)	#4 Distillation Column Condenser	6.4
32.	(H17)	#4 Distillation Column Calandria	3.7
33.	(H18)	Nitrogen Heater	1.3

TABLE 4.2-7 (Continued)

34.	(P1)	TCS Reactor Off Gas Compressor	53.2
35.	(P2)	Caustic Supply Pump	1.56
36.	(P3)	#1 Distillation Column Overheads Pump	2.64
37.	(P4)	#1 Distillation Column Calandria Pump	3.83
38.	(P5)	TET/TCS Feed Pump	2.04
39.	(P6)	#2 Distillation Column Overhead Pump	2.8
40.	(P7)	TCS Feed Pump	1.8
41.	(P8)	#2 Distillation Column Calandria Pump	3.8
42.	(P9)	#3 Distillation Column Overhead Pump	2.2
43.	(P10)	Rod Reactor TCS Feed Pump	1.7
44.	(P11)	#3 Distillation Column Calandria Pump	2.6
45.	(P12)	Rod Reactor Off Gas Compressor	235.5
46.	(P13)	#4 Distillation Column Overheads Pump	1.87
47.	(P14)	#4 Distillation Column Calandria Pump	1.87
48.	(P15)	TET Feed Pump	1.56
49.	(P16)	Waste Treatment Pump	.77
50.	(P17)	Crude TCS Feed Pump	1.9
51.	(P18)	Process Water Feed Pump	3.7
52.	(Cl)	#1 Gas Scrubber	53.2
53.	(C2)	#2 Gas Scrubber	29.
54.	(C3)	#1 Distillation Column	26.1
55.	(C4)	#2 Distillation Column	27.7
56.	(C5)	#3 Distillation Column	8.9
57.	(C6)	#4 Distillation Column	6.7
58.	(R1)	TCS Fluidized Bed Reactor	57.2
59.	(R2)	Polysilicon Rod Reactors (305)	56. (each)
60.	(Al)	Molecular Sieves	16.77
61.	(A2)	Fines Separator	4.8
62.	(A3)	Hydrogen Flare	1.
63.	(A4)	Filament Pullers (5)	15. (each)
		TOTAL PURCHASED COST	\$19,307.14 (1975 dollars) x 1.4 inflation \$27,030.00 (1980 dollars)

TABLE 4.2-8

PRODUCTION LABOR COST FOR
CONVENTIONAL POLYSILICON PROCESS

	Unit Operation	Skilled Labor Man-Hrs/Kg Si	Cost \$/kg 8i	
1.	TCS Production	.0292	.2014	
2,	Vaporization	.0219	.1511	
3.	Vapor Compression	.0219	.1511	
4.	Vapor Condensation	.0219	.1511	
5.	TCS/TET Separation	.0146	.1007	
6.	TCS Purification	.0128	.0883	
7.	TET Purification	.011	.0759	
8.	Filament Pullers	.0438	.3021	
9.	Gas Scrubbing	.0232	.1600	
10.	Hydrogen Drying (Molecular Sieves)	.0117	.0807	
11.	Crude TCS Recycle System	.0212	.1463	
12.	Silicon Fines Separation	.0055	.038	
13.	Materials Handling	.0329*	.1612*	
14.	Polysilicon Production	. 2672	1.8429	
		TOTAL COST	\$3.65 (1975 do	.1

TOTAL COST \$3.65 (1975 dollars) × 1.4 inflation 5.11 (1980 dollars)

.

^{*}semiskilled

TABLE 4.2-9
ESTIMATION OF PLANT INVESTMENT COST FOR
CONVENTIONAL POLYSILICON PROCESS

		Investment	(\$1000)
		1975 Plant	1960's Plant
1.	DIRECT PLANT INVESTMENT COSTS		
	1. Major Progess Equipment Cost	19,307	11,032
	2. Installation of Major Process Equipment	4,699	2,685
	3. Process Piping, Installed	8,969	5,125
	4. Instrumentation, Justailled	924	528
	5, Electrical, Installed	1,931	1,103
	6. Process Buildings, Installed	3,303	1,889
la.	SUBTOTAL FOR DIVECT PLANT INVESTMENT COSTS (PRIMARILY BATTERY LIMIT PACILITIES)	39,133	22,362
2.	OTHER DIRECT PLANT INVESTMENT COSTS		
	1. Utilities, Installed	9,096	5,198
	2. General Services, Site Development,		
	Fire Protection, etc.	2,317	1,324
	3. General Buildings, Offices, Shops, etc.	5,104	2,917
	4. Receiving, Shipping Facilities	4,741	2,709
2a.	SUBTOTAL FOR SHEEF FIRST PLANT INVESTMENT COSTS (PRIMARILS GUESITE FACILITIES GUESIDE BATTERY LIMITS)	21,258	12,147
3.	TOTAL DIRECT PLANT INVESTMENT COST, 14 + 2a	60,391	34,509
4.	INDIRECT PLANT INVESTMENT COSTS		
	1. Engineering, Overhead, etc.	3,757	2,147
	2. Mormal Coat, for Floods, Strikes, etc.	9,076	5,186
4a.	TOTAL INDIRECT FLANT INVESTMENT COST	12,833	7,333
5.	TOTAL DIRECT AND UNDERSION PLANT INVESTMENT COST, 3 + 1a	73,224	41,842
6.	OVERALL CONTINGENCY, 3 OF 5 010%	7,322	4,184
7.	FIXED CAPITAL INVESTMENT FOR PLANT, 5 + 6	80,546	46,026
197 196	00 CE Plant Cost Index = 253 (March) 5 CE Plant Cost Index = 182 5 CE Plant Cost Index = 104 0 CE Plant Cost Index = 102	Plant Constructed In 1975	Plant Constructed In 1960's (1965 or Earlier)

TABLE 4.2-10

ESTIMATION OF TOTAL PRODUCT COST FOR CONVENTIONAL POLYSILICON PROCESS

PRODUCT COST, \$/KG Si

		Low 1.5¢/kw hr	High 3¢/kw hr	Intermed. 2.25¢/kw hr
1.	Direct Manufacturing Cost 1. Raw Materials 2. Direct Operating Labor 3. Utilities 4. Supervision and Clerical 5. Maintenance and Repairs	15.34 3.65 8.49 .55 2.16	15.34 3.55 14.26 .55 2.16	15.34 3.65 11.37 .55 2.16
	6. Operating Supplies 7. Laboratory Charge	.55	. 55	. 55
2.	Indirect Manaufacturing Cost 1. Depraciation 2. Local Taxes 3. Insurance	.92 .46	.92 .46	.92 .46
3.	Plant Overhead	3.82	3,82	3.82
4.	By~Product Credit	(6.23)	(6.23)	(6.23)
4a.	Total Manufacturing Cost, 1 + 2 + 3 + 4	30.14	35.91	33.02
5.	General Expenses 1. Administration 2. Distribution and Sales 3. Research and Development	2.15 2.15 1.08	2.15 2.15 1.08	2.15 2.15 1.08
6.	Total Cost of Product, 4a + 5	$\frac{35.52}{\times 1.4}$	$\frac{41.29}{\times 1.4}$	38.41 (1975 dollars) x 1.4 inflation 53.16 (1980 dollars)

Basis: The above results are based on a plant constructed in 1960's (1965 or earlier) which is fully depreciated. The range reflects low and high electrical costs (1.5-3¢/kw hr). Intermediate reflects intermediate electrical cost (2.25¢/kw hr).

4.3 UCC Silane Process for Silicon (Union Carbide Corporation)

The economic analysis activity involves a cost analysis of the process under consideration for the production of silicon. The cost analysis for the particular technology is based on process design results, such as requirements for raw materials and major process equipment necessary to produce the product, from the chemical engineering analysis activity. Primary results issuing from the economic analysis include plant capital investment and product cost which are useful in identification of those processes showing promise for meeting project cost goals.

The cost analysis results for producing silicon by the UCC silane process (Union Carbide Corporation) are presented in Table 4.3-1 including costs for raw materials, labor, utilities and other items composing the product cost (total cost of producing silicon). The tabulation summarizes all of these items to give a total product cost without profit of \$6.90 (1975 dollars) and \$9.66 (1980 dollars) per kg. This product cost without profit includes direct manufacturing cost, indirect manufacturing cost, plant overhead and general expenses.

The product cost represents all cost associated with producing 1 kg of silicon. On top of these costs a producing company will include some profit. The sales price of the product silicon will actually be the sum of the product cost and a profit for the company. The profit is usally measured in terms of rate of return on the capital investment that the company spent in going into the polysilicon business. Two profitability methods which are commonly used are the return on original investment (per cent ROI) and discounted cash flow rate of return (per cent DCF).

The cost and profitability analysis summary for this process are presented in Table 4.3-2. The sales price of polysilicon at various rates of return for both profitability methods (per cent ROI and DCF) is shown in the lower half of the table. The results indicate a sales price of \$13 per kg of silicon (1980 dollars) at 15 per cent DCF return on investment.

These cost and profitability results for the UCC silane process indicate that this new technology for producing polysilicon shows good promise for meeting the cost goal of \$14 per kg of silicon material (1980 dollars) for solar cells.

The detailed results for the economic analysis are presented in a tabular format to make it easier to locate cost items of specific interest. The guide for the tabular format is given below:

	Preliminary Economic Analysis Activities Table	
•	Process Design Inputs	4.3-4
٠	Base Case Conditions	4.3-5
•	Raw Material Cost	4.3-6
٠	Utility Cost	4.3 - 7
٠	Major Process Equipment Cost	4.3-8
	Production Labor Cost	4.3-9
•	Plant Investment	4.3-10
	Total Product Cost Table	

The economic analysis provides detailed cost data for raw materials, utilities, labor and major process equipment which are necessary for polysilicon production.

TABLE 4.3-1
ESTIMATION OF PRODUCT COST FOR UCC Silane Process

			Cost of Silicon 5 dollars)		Cost of Silicon O dollars)
1.	Direct Manufacturing Cost (Direct Costs) Raw Materials Direct Operating Labor Utilities Supervision and Clerical Maintenance and Repairs Operating Supplies Laboratory Charge	•	4.15		5.81
2.	Indirect Manufactuing Cost (Fixed Cost) Depreciation Local Taxes Insurance	•	1.19		1.67
3.	Plant Overhead	•	0.66		0.92
4.	General Expenses	•	0.90		1.26
5.	Product Cost Without Profit	•	6.90	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	9,66

TABLE 4.3-2

COST AND PROFITABILITY ANALYSIS SUMMARY FOR UCC Silane Process

2. 3. 4.	Process Plant Size Plant Product Product Form Plant Investment			1,000 Metric .Silicon .Liquid Phas \$10,570,000	Tons/year
	Worl	ed Capital king Capita 15%) T	1	\$9.19 Mega \$1.38 Mega \$10.57 Mega (1975 dollars)	\$12.87Mega <u>\$ 1.99Mega</u> \$14.8(Mega (1980 dollars)

6. Return on Original Investment, after taxes (NROI)

	Sales Price \$/Kg of Silicon (1975 dollars)	Sales Price \$/Kg of Silicon (1980 dollars)
0% ROI	6.90	9.66
5% ROI	7.88	11.02
10% ROI	8.86	12,39
15% ROI	9,84	13.75
20% ROI	10.81	15,11
25% ROI	11.79	16.47
30% ROI	12.77	17.84
40% ROI	. 14.73	20.56

7. Discounted C.sh Flow Rate of Return, after taxes (* DCF)

		\$/Kg of Silicon (1975 dollars)	\$/Kg of Silicon (1980 dollars)
0%	DCF	6.90	9.66
5%	DCF	7.53	10.54
10%	DCF	8,22	11.50
15%	DCF	8.97	12.55
20%	DCF	9.77	13.65
25%	DCF	10.60	14.82
30%	DCF	11.47	16.02
	DCF		18.53

Based on 10 year project life and 10 year straight line depreciation.

8. Tax Rate (Federal) ······46%

Table 4.3-3 ECCNOMIC ANALYSES: PRELIMINARY ECONOMIC ANALYSIS ACTIVITIES FOR UCC Silane Process

	Prel. Process Economic Activity	Status		Prel. Process Economic Activity "atus
ı.	Process Design Inputs	•	6.	Production Labor Costs
	l. Raw Material Requirements	9		1. Base Cost Per Man Hour
	 Utility Requirements 	8		Cost/Kg Silicon Per Area
	3. Equipment List	9		3. Total Cost/Kg Silicon
	4. Labor Requirements	G		
			7.	Estimation of Plant Investment
2.	Specify Base Case Conditions	•		 Battery Limits Direct Costs
	 Base Year for Costs 	6		2. Other Direct Costs •
	Appropriate Indices for Costs	•		3. Indirect Costs
	 Additional 	c		4. Contingency
				5. Total Plant Investment
3.	Raw Material Costs	•		(Fixed Capital)
	 Base Cost/Lb. of Material 	e		
	Material Cost/Kg of Silicon	2	ê,	Estimation of Total Product Cost
	Total Cost/Kg of Silicon	2		1. Direct Manufacturing Cost
				2. Indirect Manufacturing Cost
4.	Utility Costs	•		3. Plant Overhead
	1. Base Cost for Each Utility	•		4. By-Product Credit •
	2. Utility Cost/Kg of Silicon	9		5. General Expenses
	3. Total Cost/Kg of Silicon	•		6. Total Cost of Product
5.	Major Process Equipment Costs	•		
	 Individual Equipment Cost 	e		
	2. Cost Index Adjustment	*		
	-			O Plan
				• In Progress
				€ Complete

TABLE 4.3-4

PROCESS DESIGN INPUTS FOR UCC SILANE PROCESS

- 1. Raw Material Requirements
 - -M.G. Silicon, milicon tetrachloride, hydrogen, copper catalyst, lime
 - -see table for "Raw Material Cost"
- 2. Utility
 - -electrical, steam, cooling water, etc.
 - -see table for "Utility Cost"
- 3. Equipment List
 - -93 pieces of major process equipment
 - -process vessels, heat exchangers, reactor, etc.
- 4. Labor Requirements
 - -production labor for purification, vaporization, product handling, etc.
 - -see table for "Production Labor Cost"

TABLE 4.3-5

BASE CASE CONDITIONS FOR UCC SILANE RPOCESS

1. Capital Equipment

```
-January 1975 Cost Index for Capital Equipment Cost
-January 1975 Cost Index Value = 430
```

2. Utilities

- -Electrical, Steam, Cooling Water, Nitrogen
- -January 1975 Cost Index (U. S. Dept. Labor)
- -Values determined by literature search and summarized in cost standardization work

3. Raw Material Cost

- -Chemical Marketing Reporter
- "January 1975 Value
- -Row Materia! Cost Index for Industrial Chemicals
- -1975 Cost Index Value = 206.9 (Wholesale Price Index, Producer Price Index)

4. Labor Cost

- -Average for Chemical Petroleum, Coal and Allied '.. Justries (1975)
- -Skilled \$6.90/hr
- -Semiskilled \$4.90/hr

5. Update to 1980

- -historically cited 1975 dollars (LSA project)
- -DOE decision to change to 1980 dollars (JPL,6/22/79)
- -reports to reflect both 1975 and 1980 dollars (JPL, 6/22/79)
- -inflation factor of 1.4 to be used (JPL, 6/22/79)

TABLE 4.3-6 RAW MATERIAL COST FOR UCC SILANE PROCESS

Raw Material	Requirement lb/KG of Si	\$/lb of Material	Cost \$/KG of Si
l. M.G. Silicon (Si)	2.60	0.535	1.391
 Silicon Tetrachloride (SiCl₄, make-up) 	2.76	0.135	0.373
 Liquid Hydrogen (H₂, make~up) 	0.032	1.84	0.059
. Copper Catalyst (Cu)	0.051	0.922	0.047
. Hydrate Lime (Ca(OH) ₂)	2.43	0.015 (33.2\$/ton)	0.036
		TOTAL,	1.906 (1975 dol x 1.4 inflation 2.668 (1980 do

2.668 (1980 dollars)

TABLE 4.3-7
UTILITY COST FOR UCC SILANE PROCESS

	Utility	Requirement/KG of Silicon	Cost of Utility	Cost \$/KG of Silicon
1.	Electricity	3.050 KW	0.0324 \$/KW-HR	0.0988
2.	Steam	172.200 lbs.	1.35 \$/klb	0.2325
3.	Cooling Water	525.000 gallons	0.09 \$/kgal	0.0473
4.	Process Water	0.0709 gallons	0.405\$/kgal	0.0001
5.	Refrigerant	968.000 Btu	10.50 \$/MBtu	0.0102
6.	Fuel	27,100.00 Btu	1.40 \$/MBtu	0.0379
			TOTAL	0.43 ()975 dollars) x 1.4 inflation 0.60 (1980 dollars)

Note:

 $k = kilo = 10^3$ $M = mega = 10^6$

TABLE 4.3-8

PURCHASED COST OF MAJOR PROCESS EQUIPMENT FOR UCC SILANE PROCESS

	Equip	ment	Purchased Cost, \$1000
1.	(D-01)	Crude TCS/STC Stripping Column	5.5
2.	(D-02)	TCS/STC Distillation Column	32.6
3.	(D-03)	DCS/TCS Distillation Column	61.6
4.	(D-04)	Silane Distillation Column	50.3
5.	(R-01)	Hydrogenation Reactor	87.6
б.	(R-)	DCS Redistribution Reactor	19.2
7.	(R-03)	TCS Redistribution Teactor	17.3
8-9.	(R-0	4) Sludge Neutralization Reactor	10.3
10.	(H-01)	Liquid H2 Vaporizer (Provided by Vendor)	
11.	(H-02)	STC Cooler	26.4
12.	(H-03)	Quench Condenser	22.6
13.	(H-04)	Recycle STC Vaporizer	3.3
14.	(H-05)	Recycle STC Superheater	35.0
15.	(H-06)	Recycle H ₂ Heater	10.8
16.	(H-07)	Stripper Reboiler	1.5
17.	(H-08)	Stripper Condenser	1.4
18.	(H-09)	TCS/STC Reboiler	8.6
10.	(* 10)	TUS/STC Condenser	44.5
20.	(H-11)	DCS/TCS Reboiler	8.2
21.	(H-12)	DCS/TCS Condenser	16.2
22.	(H-13)	DCS Cooler	1.5
23.	(H-14)	TCS Cooler	3.0

TABLE 4.3-8 (Continued)

24.	(H-15)	Silane Reboiler	1.3
25.	(H-16)	Silane Condenser	2.6
26.	(H-17)	Silane Vaporizer/Superheater	2.4
27.	(H-18)	Pyrolysis Hydrogen Cooler	4.1
28.	(H-19)	First Stage H2 Intercooler	3.6
29.	(H-20)	Second Stage H2 Intercooler	3.6
30.	(C-01)	Pneumatic Conveying Fan	1.6
31.	(C-02)	Recycle H ₂ Blower	4.7
32.	(C-03)	First Stage H ₂ Compressor	9.7
33.	(C-04)	Second Stage H ₂ Compressor	9.7
34.	(C-05)	Third Stage H ₂ Compressor	9.7
35.	(P-01)	Quench Contactor Pump	2.4
36.	(P-03)	Recycle STC Pump	15.0
37.	(P-04)	TCS Distillate Pump	19.8
38.	(P-05)	DCS Distillate Pump	11.3
39.	(P-06)	Lime Tank Pump	2.0
40.	(T-01)	Crude TCS/STC Storage Tank	39.0
41.	(T-02)	STC Storage Tank	17.0
42.	(T-03)	Liquid H ₂ Storage (Provided By Vendor)	
43.	(T-04)	Waste Settler Tank	27.0
44.	(T-05)	Waste Chloride Tank	1.8
45.	(T-06)	Quench Condenser Receiver	8.8
46.	(T-07)	Recycle H ₂ Receiver	7.2

TABLE 4.3-8 (Continued)

47.	(T-08)	Stripper Reflux pot	1.2
48.	(T-09)	TCS/STC Feflux pot	6.1
49.	(T-10)	LCS/TCS Reflux pot	11.2
50.	(T-11)	A, B Silane Shift Tank (two)	20.6 ea.
51.	(T-13)	Pyrolysis H ₂ Receiver	7.9
52.	(T-14)	Lime Make-up Tank	5.7
53.	(T-15)	Sludge Pump Tank	11.3
54.	(B-01)	M. G. Silicon Storage Hopper	12 2
55-5	6. (B~	04) Pyrolysis Dust Bin	1.7
57.	(F-01)	Crude TCS/STC Filter	0.7
58.	(F-02)	Waste Hydroxide Filter	5.0
59.	(F-03)	Pyrolysis H ₂ Filter	0,7
60.	(F-04)	M. G. Silicon Unloading Filter	1.6
61.	(5-01)	M. G. Silicon Unloading Cyclone	1.4
62.	(S-02)	Double Shell Blender	13.0
63.	(8-03)	M. G. Silicon Loak Hopper	5.8
64.	(U=01)	Quench Contactor Ejector	1.3
65.	(p=02)	Lime Tank Agitator	1.3
66.	(Fi-+iJ)	Vent Gas Combustor	6.3
67.	(D=04)	Vent Gas Ejector	1.3
68.	(R=05)	Silane Pyrolysis Reactor (six)	46.8 ea.
69.	(X-01)	Melters (six)	53.0 ea.
70.	(B-05)	Powder Hoppers (two)	14.9 ea.
71.	(X-02)	Hydrogen Cooler	4.1

TABLE 4.3-8 (Continued)

72.	(X-03)	Hydrogen Blower		2.5	
73.	(X-04)	Dust Filter		0.8	
74.	(X-J5)	Star Valve (six)		1.2	œa.
75.	(X-06)	Conveyor		8,3	
76.	(X-07)	Drum Loader	****	16.6	
					/100# 3-11-w-\
			<u>x</u>	1.4	()975 dollars) inflation
			20	174.7	(1980 dollars)

TABLE 4.3-9 PRODUCTION LABOR COST FOR UCC SILANE PROCESS

Section/ Unit Operation		Skilled Labor Man-Hrs/KG of Si	Semiskilled Labor Man-Hrs/KG of Si	Cost \$/KG of Si
1.	Hydrogenation	0.00745	0.000745	0.0879
2.	Silane	0.02230	deal mark deal gray good damp area	0.1539
з.	Pyrolysis	0.02980	them space thank hope high talks the	0.2056
4.	Waste Treatment	0.00745	ting the stat duri and and ting the	0.0514
5.	Hydrogen Compression	0.00745		0.0514
			TOTAL	0.55 (1975 dollars) x 1.4 inclation 0.77 (1980 dollars)

TABLE 4.3-10

ESTIMATION OF PLANT INVESTMENT FOR UCC SILANE PROCESS

		Investmen \$1000	nt
1.	DIRECT PLANT INVESTMENT COSTS 1. Major Process Equipment Cost 2. Installation of Major Process Equipment 3. Process Piping, Installed 4. Instrumentation, Installed 5. Electrical, Installed 6. Process Buildings, Installed	1,481.9 637.2 1,096.6 281.6 148.2 148.2	
la.	SUBTOTAL FOR DIRECT PLANT INVESTMENT COSTS (PRIMARILY BATTERY LIMIT FACILITIES)	3,793.7	
2.	OTHER DIRECT PLANT INVESTMENT COSTS 1. Utilities, Installed 2. General Services, Site Development, Fire Protection, etc. 3. General Buildings, Offices, Shops, etc. 4. Receiving, Shipping Facilities	711.3 177.8 207.5 311.2	
2a.	SUBTOTAL FOR OTHER DIRECT PLANT INVESTMENT COSTS (PRIMARILY OFFSITE FACILITIES OUTSIDE BATTERY LIMITS)	1,407.8	
3.	TOTAL DIRECT PLANT INVESTMENT COST, la + 2a	5,201.5	
4.	 INDIRECT PLANT INVESTMENT COSTS Engineering, Overhead, etc. Normal Cont. for Floods, Strikes, etc. 	815.0 1,052.1	
4a.	TOTAL INDIRECT PLANT INVESTMENT COST	1,867.1	
5.	TOTAL DIRECT AND INDIRECT PLANT INVESTMENT COST, 3 + 4a	7,068.6	
6.	OVERALL CONTINGENCY, % of 5	2,120.6	<u>-</u>
7.	FIXED CAPITAL INVESTMENT FOR PLANT, 5 + 6	\times 1.4	(1975 dollars) inflation (1980 dollars)

TABLE 4.3-11

ESTIMATION OF TOTAL PRODUCT COST POR UCC SILANE PROCESS

		\$/KG of Bi
1.	Direct Manufacturing Cost (Direct Charges)	
	1. Raw Materials	3.906
	2. Direct Operating Labor	0.550
	3. Utilities	0.427
	4. Supervision and Clerical	0.083
	5. Maintenance and Ropairs	0.919
	6. Operating Supplies	0.184
	7. Laboratory Charge	0 083
2.	Indirect Manufacturing Cost (Fixed Charges)	
	1. Depreciation	J.919
	2. Local Taxes	0.184
	3. Insurance	0.092
3.	Plant Overhead	0.656
4.	By-Product Credit	No. on on on
4a.	Total Manufacturing Cost, 1 + 2 + 3 + 4	6 003
5.	General Expenses	
	1. Administration	0.360
	2. Distribution and Sales	0.360
	3. Research and Development	0 180
6.	Total Cost of Product, 4a + 5	6.903 (1975 dollars)
٠.	Controvat of Floring Cy 10 1 3	x 4 inflation
		9 664 (1980 dollars)

4.4 BCL Process for Silicon - Case A (Battelle Columbus Laboratories)

The economic analysis activity involves a cost analysis of the process under consideration for the production of silicon. The cost analysis for the particular technology is based on process design results, such as requirements for raw materials and major process equipment necessary to produce the product, from the chemical engineering analysis activity. Primary results issuing from the economic analysis include plant capital investment and product cost which are useful in identification of those processes showing promise for meeting project cost goals.

The cost analysis results for producing silicon by the BCL process-Case A (Battelle Columbus Laboratories) are presented in Table 4.4-1 including costs for raw materials, labor, utilities and other items composing the product cost (total cost of producing silicon). The tabulation summarizes all of these items to give a total product cost without profit of \$8.63 (1975 dollars) and \$12.08 (1980 dollars) per kg. This product cost without profit includes direct manufacturing cost, indirect manufacturing cost, plant overhead and general expenses.

The product cost represents all cost associated with producing silicon. On top of these costs a producing company will include some profit. The sales price of the product silicon will actually be the sum of the product cost and a profit for the company. The profit is usually measured in terms of rate of return on the capital investment that the company spent in going into the polysilicon business. Two profitability methods which are commonly used are the return on original investment (per cent ROI) and discounted cash flow rate of return (per cent DCF).

The cost and profitability analysis summary for this process are presented in Table 4.4-2. The sales price of polysilicon at various rates of return for both profitability methods (per cent ROI and DCF) is shown in the lower half of the table. The results indicate a sales price of \$13.28 per kg of silicon (1980 dollars) at 5 per cent DCF return on investment after taxes.

The detailed results for the economic analysis are presented in a tabular format to make it easier to locate cost items of specific interest. The guide for the tabular format is given below:

•	Preliminary Economic Analysis Activities Table	4.4 - 3
•	Process Design Inputs	4.4-4
•	Base Case Conditions	4.4 - 5
•	Raw Material Cost	4.4 - 6
•	Utility CostTable	4.4 - 7
٠	Major Process Equipment Cost	4.4-8
	Production Labor Cost	
•	Plant Investment	4.4 - 10
•	Total Product Cost	4.4 - 11

These cost and profitability results for the BCL process-Case A indicate that this new technology for producing polysilicon shows promise for meeting the cost goal of \$14 per kg of silicon material (1980 dollars) for solar cells.

TAPLE 4.4-1
ESTIMATION OF PRODUCT COST FOR BCL PROCESS - CASE A

		Cost \$/Kg of Silicon (1975 dollars)	Cost \$/Kg of Silicon (1980 dollars)
1.	Direct Manufacturing Cost (Direct Costs) Raw Materials Direct Operating Labor Utilities Supervision and Clerical Maintenance and Repairs Operating Supplies Laboratory Charge	. 5.21	7.29
2.	Indirect Manufactuing Cost (Fixed Cost) Depreciation Local Taxes Insurance	. 1.62	2.27
3.	Plant Overhead	. 0.68 '	0.95
4.	General Expenses	. 1.12	1.57
5.	Product Cost Without Profit	. 8.63	12.08

TABLE 4.4-2

COST AND PROFITABILITY ANALYSIS SUMMARY FOR BCL PROCESS-CASE A

	Process		ase A Tons/year
	Plant Product.		
4.	Product Form	····Silicon Granu	les
5.	Plant Investment	(1975 dollars)	\$20,070,000 (1980 dollars)
	Fixed Capital Working Capital	\$12.47Mega \$ 1.87Mege	\$17.45 Mega \$ 2.62 Mega
	(15%) Total	\$14.34Mega	\$20.07 Maga
	(154) 10051	(1975 dollars)	(1980 dollars)

6. Return on Original Investment, after taxes (%ROI)

		Sales Pricq \$/Kg of Silicon (1975 dollars)	Sales Price \$/Kg of Silicon (1980 dollars)
0%	ROI.	8.63	13.08
51	ROI	9.96	13.94
10%	ROI	11,28	15.80
15%	ROI	12.61	17.65
20%	ROI	13,94	19.51
	ROT		21.37
	ROI	1 / P	23.23
	ROI	19.25	26 95

7. Discounted Cash Flow Rate of Return, after taxes (t DCF)

		Sales Price \$/Kg of Silicon (1975 dollars)	Saies Price S/Kg of Sillicon (1980 dolla: 4)
OR	DCF	8.63	12.78
54	DCF	9.48	13 28
10%	DCF	10.42	34 5∋
154	DCF	11.44	16 Ca
20%	DCT	12.52	17 b₃
254	PCF	13.65	te 11
30%	DCF	14.83	233 755
40%	DCF	17.27	24 73

Based on 10 year project life and $10^{-10000}~\text{straight}$ line depreciation.

8. Tax Rate (Federal).....

TABLE 4.4-3

ECONOMIC ANALYSES: PRELIMINARY ECONOMIC ANALYSIS ACTIVITIES FOR BCL PROCESS-CASE A

	Prel. Process Enchants Astracty	Status	Prel. Process Economic Activity	Status
-	Process Design Inputs	•	1. Frowlation Lazor Costs	•
	i, Raw Material Peq 1 ordnis	•	1. Base Cost Per Man Hour	•
	3. Obility Regulation	•	1. Josh Yg Erfloom Ber Area	•
	D Equipment Cust	7	3. Juai Chat/Mg 2:1./ xx ;	•
	4. Tabor Brightzerkin			
			ascumation of Trant Tromschent	•
	pen fy sask (1986-1987)	•	1. Battery Limits Direct Osts	•
	. Same four for	•	1. Tope: Lireut Costs	•
	App op plate Indio Color	•	3. unditent Costs	
	Fig. Lines	•	4 year open by	•
			5. ocal Plant Investment	•
7	Pain Kutheru e	2	: ked lagut al ,	
	The Base of the Arms of	•		
	factor in the	•	Electration of Total Troduct Cost	•
	Constant was the second	7	1. lireyo Manufastaring Cost	•
	•		<pre>1. indirect Manufacturing Cost</pre>	•
7.	Top Att Comment	7	J Plant Overhead	6
	1 Bark Corp For La 1197y	-	By-Product Oredut	•
	- 2. Calify in Wigh Sales A	•	J. General Expenses	•
	3. Motal Distifug of Course		4. Total Cost of Product	•
5.	Majok Trocess Equipment Costs	÷		
_	1. Individual Equipment Cost	*		
	2. Cost Index Adjustment	•		
			3 Plan	
			€ In Progress	
			• Complete	

TABLE 4.4-4

PROCESS DESIGN INPUTS FOR BCL PROCESS - Case A

- 1. Raw Material Requirements
 -Silicon tetrachloride, zinc, lime, argon and nitrogen
 -see table for "Raw Material Cost"
- 2. Utility -electricity, steam, cooling water and process water -see table for "Utility Cost"
- 3. Equipment List
 -82 plus pieces of major process equipment
 -process vessels, heat exchangers, reactor, etc.
- 4. Labor Requirements -production labor for purification, deposition, electrolysis, etc. -see table for "Production Labor Cost"

TABLE 4.4-5 BASE CASE CONDITION FOR BCL PROCESS-CASE A

1. Capital Equipment

- -January 1975 Cost Index for Capital Equipment Cost
- -January 1975 Cost Index Value * 430

2. Utilities

- -Electrical, Steam, Cooling Water, Nitrogen
- -January 1975 Cost Index (U. S. Dept. Labor)
- "Values determined by literature search and summarized in cost standardization work

3. Raw Material Cost

- -Chemical Marketing Reporter
- -January 1975 Value
- -Raw Material Cost Index for Industrial Chemicals
- -1975 Cost Index Malue (2000) Wholesale Price Index, Producer Price Index)

4. Labor Cost

- -Average for Chemical Petroleum, Coal and Allied Industries (1975) -Skilled \$6.907h;
- 5. Update to 1980
 - -historically cited 1975 dollars (ISA project)
 - -DOE decision to change to 1980 dellars (JPL, 6/22/79)
 - -reports to reflect both 1975 and 1980 dollars (JPL, 6/22/79)
 - -inflation factor of 1.4 to be used (JPL, 6/22/79)

Table 4.4-6

RAW MATERIAL COST FOR BCL PROCESS-CASE A

	Raw Material	Requirement 1b/KG of Si	\$/1b of Material	Cost \$/KG of Si
1.	Silicon Tetrachloride (SiCl ₄)	15.68	0.135	2.117
2.	Zinc (Zn)	0.54	0.38	0.205
3.	Hydrate Lime (Ca(OH) ₂)	2.85	0.015	0.043
4.	Argon (Ar)	3.1 SCF	0.016/scF	0.050
5.	Nitrogen (N2)	7.6 SCF	0.003/scf	0.023
			Sub Total	2.438
6.	Chlorine (Cl ₂)	-10,46 ¹	0.0332	-0.347
			TOTAL	2.091 (1975 dollars) x 1.4 inflation 2.927 (1980 dollars)

Note:

^{1.} This number is the result of by-product rate minus reactor chlorination rate, i.e., 11.12 - 0.66 lb. of Cl_2/KG Si.

Table 4.4-7
UTILITY COST FOR BCL PROCESS-CASE A

	Utility	Requirement/KG of Silicon	Cost of Utility	Cost \$/KG of Silicon
1.	Electricity	30.92 kw-hr	0.0324 \$/kw-hr	1.0018
2.	Steat.	9.67 pounds	1.35 \$/klb	0.0131
3.	Cooling Water	37.88 Gallons	0.09 \$/kgal	0.0034
4.	Process Water	24.20 Gallons	0.405 \$/kgal	0.0098
5.	Refrigerant	2.38 MBtu	10.50 \$/MBtu	0.0250
			TOTAL	1.0531 (1975 dollars) x 1.4 inflation 1.4743 (1980 dollars)

Note:

 $k = kilo = 10^3$ $M = mega = 10^6$

TABLE 4.4-8
ESTIMATED COST OF MAJOR PROCESS EQUIPMENT FOR BCL PROCESS-CASE A

Equipment		Equipment	Purchased Cost, \$1,000	
1.	(D-01)	Light End Distillation Column	55.6	
2.	(D-02)	Heavy End Distillation Column	55.6	
3.	(A-01)	Primary SiCl ₄ Vent Scrubber	0.8	
4.	(A-02)	Final SiCl ₄ Vent Scrubber	11.1	
5.	(H-01)	L. E. Column Feed Heater	7.8	
6.	(H-02)	L. E. Column Reboiler	2.2	
7.	(H-03)	L. E. Column Condenser	2.3	
8.	(H-04)	H. E. Column Feed Heater	7.8	
9.	(H-05)	H. E. Column Reboiler	2.4	
10.	(H-06)	H. E. Column Condenser	2.3	
11.	(H-07)	SiCl ₄ Vent Condenser	11.8	
12.	(H-08)	SiCl ₄ Vaporizer	6.7	
13.	(H-09)	Reactor Condensers (2)	190.3	
14.	(H-10)	Reactor ZuCl ₂ Strippers (2)	27.9	
15.	(H-11)	SiCl ₄ Condenser	20.5	
16.	(H-12)	Cell ZnCl ₂ Stripper	10.9	
17.	(H-13)	Therminol Cooler (Cold Circuit)	3.8	
18.	(H-14)	Therminol Cooler (Hot Circuit)	9.1	
19.	(H-15)	Start-up Heater	9.6	
20.	(H-16)	Silicon Product Coolers (2)	7.7	
20a.	(H-17)	Chlorination Cooler	15.9	
20b.	(H~18)	Cell Gas Cooler	18.7	
21.	(T-01)	SiCl ₄ Storage Tank	33.6	
22.	(T-02)	SiCl ₄ Emergency Storage Tank	33.6	
23.	(T-03)	L. E. Column Reflux Drum	6.7	

TABLE 4.4-8 (Continued)

24.	(T-04)	Surge Tank	19.0
25.	(T-05)	Sump Tank	19.0
26.	(T-06)	H. E. Column Reflux Drum	6.7
27.	(T-07)	Pure SiCl ₄ Storage Tank	28.8
28.	(T-08)	Electrolysis Feed Tank	46.0
29.	(T-09)	Molten Zinc Storage Tank	86.9
30.	(T-10)	Therminol Head Tank	3.8
31.	(T-11)	Therminol Drain Down Tank	5.3
32.	(T-12)	Chlorine Supply Tank	2.4
33.	(T-13)	Lime Solution Storage Tank	6.8
34.	(P-01)	Purification Feed Pump	3.7
35.	(P-02)	L. E. Column Feed Pump	9.4
36.	(P-03)	L. E. Column Reflux Pump	8.4
37.	(P-04)	Surge Tank Pump	9.8
38.	(P-05)	Sump Pump	3.7
39.	(P-06)	L. E. Column Bottom Pump	12.0
40.	(P-07)	H. E. Column Reflux Pump	8.4
41.	(P-08)	H. E. Column Bottom Pump	10.9
42.	(P-09)	SiCl ₄ Vaporizer Feed Pump	4.8
43.	(P-10)	Reactor Condenser Circulation Pumps (2)	14.4
44.	(P-11)	Cold Circuit Pump	6.7
45.	(P-12)	Hot Circuit Pump	13.9
46.	(P-13)	Primary Scrubber Recirculation Pump	0.9
47.	(P-14)	Primary Scrubber Lower-loop Recirculation Pump	1.4
48.	(P-15)	Primary Scrubber Upper-loop Recirculation Pump	1.5
AΩ	(p=16)	Time Solution Metaring Dumn	1.4

TABLE 4.4-8 (Continued)

50.	(F-Ol) L. E. Column Feed Filter		0.9
51.	(F-02) L. E. Column Reflux Filter		0.9
52.	(F-03) H. E. Column Feed Filter		0.9
53.	(F-04) H. E. Column Reflux Filter		0.9
54.	(F-O5) Therminol Cooler Blower Filter		0.7
55.	(R-Ol) Fludized Bed Reactors (2)		197.1
56.	(FN-01) Furnaces (2)		354.2
57.	(B-01) Seed Addition Hoppers (2)		9.6
58.	(B-02) Si Product Hoppers (4)		14.4
59.	(B-03) Zinc Hopper		2.4
60.	(C-Ol) Therminol Cooler Blower		4.8
61.	(C-02) Scrubber Vent Blower		5.4
62.	(E-Ol) Eductors (2)		1,3
63.	(EC-01) Electrolysis Cells (6)		441.
64.	(PW-01) Fower Supply and Bus		405.9
65.	(VP=01) Zinc Vaporizers (2)		144.0
		TOTAL	2,177,7 (1977 decr) as x 1,4 int) (t or 3,048,8 (1968 as a)
			A Maria a service of the control of

TABLE 4.4-9
PRODUCTION LABOR COST FOR BCL PROCESS-CASE A

	Section	Labor Man-Hr/Kg Si	Labor Cost \$/Man-Hr	Cost \$/Kg Si
1.	Purification	0.01402	6.90	0.0968
2.	Deposition	0,01402	6.90	0.0968
3.	Electrolysis	0.02103	6.90	0.1451
4.	Waste Treatment	0.00701	6.90	0.0484
5.	Product Handling	0.00701	6.90	0.0484
			TOTAL	0.4355 (1975 dollars) x 1.4 inflation 0.6097 (1980 dollars)

Note: Costs are 1975 Dollars

TABLE 4.4-10
ESTIMATION OF PLANT INVESTMENT FOR BCL PROCESS-CASE A

		Investment \$1000
1.	DIRECT PLANT INVESTMENT COSTS	
	1. Major Process Equipment Cost	2,177.7
	2. Installation of Major Process Equipment	936.4
	3. Process Esping, Installed	1,611.6
	4. Instrumentation, Installed	413.8
	5. Electrical, Installed	217.8
	6. Process Buildings, Installed	217.8
la.	SUBTOTAL FOR DIRECT PLANT INVESTMENT COSTS (PRIMARILY BATTERY LIMIT FACILITIES)	5,574,9
2.	OTHER DIRECT PLANT INVESTMENT COSTS	
	1. Utilities, Installed	174 C. A.
	2. General Service, Site Development,	with a first
	Fire Protection, etc.	
	3. General Buildings, Offices, Chops, etc.	304.11
	4. Receiving, Shipping Facilities	457.3
2a.	SUBTOTAL FOR OTHER DIRECT PLANT INVESTMENT COSTS (PRIMARILY OFFSITE FACILITIES OUTSIDE BATTERY LIN	
3.	TOTAL DIRECT FLANT INVESTMENT COST, la + 2a	7,643.7
4.	INDIRECT PLANT INVESTMENT COSTS	
	1. Engineering, everhead, etc.	1,197.7
	2. Normal Cont. for Floods, Strikes, etc.	1,546,7
4a.	TOTAL INDIRECT DEANT INVESTMENT COST	2,74
5,	TOTAL PERSON AND INDIFFER PLANT INVESTMENT COST, 3 + 4a	19,387.6
6.	OVERTIA CONTINGENCY, * of 5	2,07.,6
7	C INVICTMENT FOR PLANT, 5 + 6	7.465. (1975 one one x 1.4 inflation 17,451.2 (1980 doctors)

TABLE 4.4-11
ESTIMATION OF TOTAL PRODUCT COST FOR BCL PROCESS- WASE A

		\$/KG of Si
1.	Direct Manufacturing Cost (Direct Charges)	
	1. Raw Materials	2,091
	2. Direct Operating Labor	0,436
	3. Utilities	1.053
	4. Supervision and Clerical	0.065
	5. Maintenance and Repairs	1.247
	6. Operating Supplies	0.249
	7. Laboratory Charge	0.065
2.	Indirect Manaufacturing Cost (Fixed Charges)	
	1. Depreciation	1,247
	2. Local Taxes	0.249
	3. Insurance	0,125
	ar anomeniae	0.14.
3.	Plant Overhead	0.675
4.	By-Product Credit	wa 100 500 507 507
Δa.	Total Manufacturing Cost, 1 + 2 + 3 + 4	7,501
5.	General Expenses	
	1. Administration	0.450
	2. Distribution and Sales	0.450
	3. Research and Development	0,225
	· · · · · · · · · · · · · · · · · · ·	Bridge and the right
ñ.	Total Cost of Product, 4a + 5	8.626 (1975 dollars)
• •	The same of the sa	x 1.4 inflation
		12,076 (1980 dollars)
		241010 (1200 Walta Di

4.5 BCL Process for Silicon - Case B (Battelle Columbus Laboratories)

The economic analysis activity involves a cost analysis of the process under consideration for the production of silicon. The cost analysis for the particular technology is based on process design results, such as requirements for raw materials and major process equipment necessary to produce the product, from the chemical engineering analysis activity. Primary results issuing from the economic analysis include plant capital investment and product cost which are useful in identification of those processes showing promise for meeting project cost goals.

The cost analysis results for producing silicon by the BCL process - Case B (Battelle Columbus Laboratories) are presented in Table 4.5-1 including costs for raw materials, labor, utilities and other items composing the product cost (total cost of producing silicon). The tabulation summarizes all of these items to give a total product cost without profit of \$7.91 (1975 dollars) and \$11.07 (1980 dollars) per kg. This product cost without profit includes direct manufacturing cost, indirect manufacturing cost, plant overhead and general expenses.

The product cost represents all cost associated with producing silicon. On top of these costs a producing company will include some profit. The sales price of the product silicon will actually be the sum of the product cost and a profit for the company. The profit is usually measured in terms of rate of return on the capital investment that the company spent in going into the polysilicon business. Two profitability methods which are commonly used are the return on original estment (per cent ROI) and discounted cash flow rate of return (per cent DCF).

The cost and profitability analysis summary for this process are presented in Table 4.5-2. The sales price of polysilicon at various rates of return for both profitability methods (per cent ROI and DCF) is shown in the lower half of the table. The results indicate a sales price of \$13.14 per kg of silicon (1980 dollars) at 10 per cent DCF return on investment.

These cost and profitability results for the BCL process-Case B indicate that this new technology for producing polysilicon shows promise for meeting the cost goal of \$14 per kg of silicon material (1980 dollars) for solar cells.

The detailed results for the economic analysis are presented in a Labular format to make it easier to locate cost items of specific interst. The guide for the tabular format is given below:

	Preliminary Economic Analysis Activities Table	4.5 - 3
٠	Process Design InputsTable	4.5 - 4
•	Base Case Conditions	4.5 - 5
	Raw Material Cost	
	Utility CostTable	
٠	Major Process Equipment Cost	4.5-8
	Production Labor Cost	
•	Plant Investment	4.5 - 10
	Total Product Cost	4.5 - 11

TABLE 4.5-1

EBTIMATION OF PRODUCT COST FOR BCL PROCESS - CASE B

		Cost \$/Kg of Silicon (1975 dollars)	Cost \$/Kq of Silicon (1980 dollars)
1.	Direct Manufacturing Cost (Direct Costs) Raw Materials Direct Operating Tabor Utilities Supervision and Clerical Maintenance and Repairs Operating Supplies Laboratory Charge	4.94	6, 52
2,	Indiract Manufactuing Cost (Fixed Cost) Depreciation Local Taxes Insurance	1.33	1,86
3.	Plant Overhead	61	, 85
4.	General Expenses	1.0	1.44
5.	Product Cost Without Profit	7.91	(1.0)

TABLE 4.5-2

COST AND PROFITABILITY ANALYSIS SUMMARY FOR BCL PROCESS- CASE B

1. 2. 3. 4.	Process Plant Size Plant Product Product Form	1,000 Metric 'SiliconSilicon Granu	Tons/vear les
5.	Plant Investment	(1975 dollars)	\$16,500,000 (1980 dollars)
	Fixed Capital Working Capital (15%) Total	\$10.25Mega <u>\$ 1.54Mega</u> \$11.79Mega (1975 dollars)	\$14.35Mega \$-2.15Mega \$15.55Mega (1980 dol:ars)

6. Return on Original Investment, after taxes (%ROI)

		Sales Price \$/Kg of Silicon (1975 dollars)	Sales Price \$/Kg of Silicor. (1980 dollars)
0%	ROI.,	7.91	11.07
50	RO1	9.00	12.60
10%	RO1	∦(1, O\)	14 13
15%	ROI	11.148	15.55
	ROI		1000
	ROI		18.71
	ROI	14 46	20 . 4
	ROI	16.64	73, 29

7. Discounted Cash Flow Rate of Return, after taxes (* DCF)

		Sales Price \$/Kg of Silicon (1975 dollars)	Sales Price \$/Kg of Siltion (1460 do 11/s)
04	DCF	7.01	¥1.0°
5%	DCF	192 to 1	12.05
10%	DCF	9,39	13.14
15%	DCF	10.72	14,34
20%	DCF	11.11	42 33
25%	DCF	12.04	16 35
	DCF	1 3 11	18.20
-	DCT		21 cm.

Based on 10 year project life and 10 year straight line depreciation.

8. Tax Rate (Federal).....40%

TABLE 4.5-3

ECONOMIC ANALYSES: PRELIMINARY ETONOMIC ANALYSIS ACTIVITIES FOR BCL PROCESS-CASE B

1. Process Design Inputs		8. Production Labor Costs	4
1. Raw Material Pequirements	•	1. Base Jost Per Man Hour	
2. Utility Requirements		1. Cost/Kg Silicon Per Area	•
i. Equipment List	<u>.</u>	3. Total Cost/Kg Silicon	•
4. Labor Requirements	•	Ji IJ GA GUACANG UALAGAN	•
the same of the desirence of	-	Es ination of Plant Investment	4
Specify Base Case Conditions	a	1. attery Limits Direct Costs	
1. Base Year for Dools	ŧ	1. ther Direct Costs	•
1 Appropriate Indices for Justs	•	indirect Costs	
Additional	•	4. Contingency	•
		5 Total Plant Investment	•
3. Pak laterial Costs	¥	Tuxud Capital)	
 Base Cost Lb. of Material 	•		
banemiel (r ist K y of Sil loor	•	Emilian with of Total Product Cost	4
Total (bet/ Kg c Sil icon	9	1. Free t Manufacturing Cost	•
		it rest Manufacturing Cost	•
4. Thisty Costs	•	3. Pwant Overhead	•
in base cost for East Cullity	£	4. Sy - Sodact Credit	•
1 till tas/Kg Ellist	•	æ eral Expenses	- 1
3. tal Dist/Kg o Flicon	•	Total Cost of Product	•
5. Major Process Equipment Cost.	4		
 I dividual Equipment Cost 	•		
Cost Index Adjustment	•		
		. Plan	
		9 In Progress	
		● Complete	

TABLE 4.5-4

PROCESS DESIGN INPUTS FOR BCL PROCESS - CASE B

- l. Raw Material Requirements
 -Silicon tetrachloride, zinc, lime, argon and nitrogen
 -see table for "Raw Material Cost"
- 2. Utility -electricity, steam, cooling water and process water -see table for "Utility Cost"
- 3. Equipment List
 -70 plus pieces of major process equipment
 -process vessels, heat exchangers, reactor, etc.
- 4. Labor Requirements -production labor for purification, deposition, electrolysis, etc. -see table for "Production Labor Cost"

TABLE 4.5-5 BASE CASE CONDITION FOR BCL PROCESS - CASE B

1. Capital Equipment

- -January 1975 Cost Index for Capital Equipment Cost
- -January 1975 Cost Index Value = 430

2. Utilities

- -Electrical, Steam, Cooling Water, Nitrogen
- -January 1975 Cost Index (U. S. Dept. Labor)
- -Values determined by literature search and summarized in cost standardization work

3. Raw Material Cost

- -Chemical Marketing Reporter
- -January 1975 Value
- -Raw Material Cost Index for Industrial Chemicals
- -1975 Cost Index Value = 206.9 (Wholesale Price Index, Producer Price Index)

4. Labor Cost

- -Average for Chemical Petroleum, Coal and Allied Industries (1975) -Skilled \$6.90/hr
- 5. Update to 1980
 - -historically cited 1975 dollars (LSA project)
 - -DOE decision to change to 1980 dollars (JPL, 6/22/79)
 - -reports to reflect both 1975 and 1980 dollars (JPL, 6/22/79)
 - -inflation factor of 1.4 to be used (JPL, 6/22/79)

Table 4.5-6

RAW MATERIAL COST FOR BCL PROCESS-CASE B

	Raw Material	Requirement lb/KG of Si	\$/lb of Material	Cost \$/KG of Si
1.	Silicon Tetrachloride (SiCl ₄)	15.68	0.135	2.117
2.	Zinc (Zn)	0.54	0,38	0.205
3.	Hydrate Lime (Ca(OH) ₂)	2.85	0.015	0.043
4.	Argon (Ar)	3.1 scf	0.016/SCF	0.050
r,	Nitrogen (Np)	7.6 821	0.003/scF	0.023
			Sub Total	2.438
6.	Chlorine (Cl ₂)	-10.461	0.032	<u>-0.347</u>
			TOTAL.	2.091 (1975 dollars) x 1.4 inflation 2.927 (1980 dollars)

Note:

^{1.} This number is the result of by-product rate minus reactor chlorination rate, i.e., 11.12 ± 0.66 Hz, of Cl_2/KG Si.

Table 4.5-7
UTILITY COST FOR BCL PROCESS-CASE B

	Utility	Requirement/KG of Silicon	Cost of Utility	Cost \$/KG of Silicon
ı.	Electricity	30.92 KW-PR	0.0324 \$/KW-HR	1.0018
2.	Steam	9.67 pounds	1.35 \$/Mlb	0.0131
3,	Cooling Water	37.88 Gallons	0.09 \$/Mgal	0.0034
4.	Process Water	24.20 Gallons	0.405 \$/Mgal	0.0098
5.	Refrigerant	2.38 MBtu	10.50 \$/MMBtu	0.0250
			TOTAL	1.0531 (1975 dollars) x 1.4 inflation 1.4743 (1980 dollars)

TABLE 4.5-8
ESTIMATED COST OF HAJOR PROCESS EQUIPMENT FOR BCL PROCESS-CASE B

	Equipment	Purchased Cost, \$1,000
1.	(D-01) Light End Distillation Column	55.6
2.	(D-02) Heavy End Distillation Column	55.6
3.	(A-O1) Primary SiCl ₄ Vent Schubber	0.8
4.	(A-02) Final SiCl ₄ Vent Scrubber	11.1
5.	(H-01) L. E. Column Feed Heater	7.8
6.	(H-O2) L. F. William Reboiler	2,2
7.	(H-05) L. F. Column Condenser	2.3
8.	(H-04) H. E. Column Feed Heater	7.8
9.	(H-05) H. E. Column Reboiler	2.4
10.	(H-06) H. E. Column Condenser	2.3
11.	(H-07) SiCi4 Vent Condenser	11.8
12.	(H-08) SiCl ₄ Vaporizer	6.7
13.	(H-09) Reactor Condensers (2)	144.2
14.	(H-10) Reactor EnCl ₂ Strippers (2)	21.1
15.	(H-11) SiCl ₄ Condenser	20.5
16.	(H-12) Cell Uncly Stripper	10.9
17.	(H-13) Therminol Cooler (Cold Circuit)	3.8
18.	(H-14) Therminol Cooler (Hot Circuit)	9.1
19.	(H-15) Start-up Heater	9.6
20.	(H-16) Silicon Product Coolers (2)	5,8
20a.	(H-1') Chlarination Cooler	15.9
20b.	(H-18) Cell Gas Cooler	18.7
21.	(T-01) Sicl ₄ Storage Tank	33.6
22.	(T-02) SiC 4 Emergency Storage Tank	33.6
23.	(T-03) I. F. Column Reflux Drum	6.7

TABLE 4.5-P (Continued)

24.	(T-04)	Surge Tank	19.0
25.	(T-05)	Sump Tank	19.0
26.	(T-06)	H. E. Column Reflux Drum.	6.7
27.	(T-07)	Pure SiCl ₄ Storage Tank	28.8
28.	(T08)	Electrolysis Feed Tank	46.0
29.	(T-09)	Molten Zinc Storage Tank	86.9
30.	(T-10)	Therminol Head Tank	3.8
31.	(T-11)	Therminol Drain Down Tank	5.3
32.	(T-12)	Chlorine Supply Tank	2.4
33.	(T-13)	Lime Solution Storage Tank	6.8
34.	(P-01)	Purification Feed Pump	3.7
35.	(P-02)	L. E. Column Feed Pump	8.4
36.	(P-03)	L. E. Column Reflux Prmy	8.4
37.	(P-04)	Surge Tank Pump	9.8
38.	(P-05)	Sump Pump	3.7
39.	(P=06)	L. E. Column Bottom Pump	12.0
40.	(P-07)	H. E. Column Reflux Pump	8.4
41.	(P-08)	H. E. Column Pottom Pump	10.9
42.	(P-09)	SiCl ₄ Vaporizer Feed Pump	4.8
43.	(P-10)	Reactor Condenser Circulation bumps (2)	10.9
44.	(P-11)	Cold Circuit Pump	6.7
45.	(P-12)	Hot Circuit Pump	13.9
46.	(P-13)	Primary Scrubber Recirculation Pump	0.9
47.	(P-14)	Primary Scrubber Lower-loop Recirculation Pump	1.4
48.	(P-15)	Primary Scrubber Upper-loop Recirculation Pump	1.5
49.	(P-16)	Lime Solution Metering Pump	1.4

TABLE 4.5-8 (Continued)

50.	(F-01) L. E. Column Feed Filter		0.9
51.	(F-02) L. E. Column Reflux Filter		0.9
52.	(F-03) H. E. Column Feed Filter		0.9
53.	(F-04) H. E. Column Reflux Filter		9.9
54.	(F-05) Therminol Cooler Blower Filter		0.7
55.	(R-Ol) Fludized Bed Reactors		149.4
56.	(FN-01) Furnaces		268.5
57.	(B-01) Seed Addition Hoppers		7.3
58.	(B-02) Si Product Hoppers (4)		14.4
59.	(B-03) Zinc Hopper		2.4
60.	(C-Ol) Therminol Cooler Blower		4.8
61.	(C-02) Scrubber Vent Blower		5.4
62.	(E-01) Eductors (2)		1.3
63.	(EC-01) Electrolysis Culls (2)		286.5
64.	(PW-01) Power Supply and Bus		105.9
65.	(VP-01) Zinc Vaporizers		109.1
		TOTAL	1,790.7 (1975 dollars) x 1.4 inflation 2,507.0 (1980 dollars)

TAB: 4.5-9
PRODUCTION LABOR COST FOR BCL PROCESS-CASE B

	Section	Labor Man-Hr/Kg Si	Labor Cost \$/Man-Hr	Cost \$/Kg Si
1.	Purification	0.01402	6.90	0.0968
2.	Deposition	0.01402	6.90	0.0968
3.	Electrolysis	0.02103	6.90	0.1451
4.	Waste Treatment	0.00701	6.90	0.0484
5.	Product Handling	0.00701	6.90	0.0484
			TOTAL.	0.4355 (1975 dollars) x 1.4 inflation 0.6097 (1980 dollars)

Note: Costs are 1975 Dollars

TABLE 4.5-10
ESTIMATION OF PLANT INVESTMENT FOR BCL PROCESS-CASE B

		Investment \$1000	: -
1.	DIRECT PLANT INVESTMENT COSTS 1. Major Process Equipment Cost 2. Installation of Major Process Equipment 3. Process Piping, Installed	1,790.7 770.0 1,325.1	
	 Instrumentation, Installed Electrical, Installed Process Buildings, Installed 	340.2 179.1 179.1	
la.	SUBTOTAL FOR DIRECT PLANT INVESTMENT COSTS (PRIMARILY PATTERY LIMIT FACILITIES)	4,584.2	
2.	OTHER DIRECT PLANT INVESTMENT COSTS 1. Utilities, Installed 2. General Service, Site Development, Fire Protection, etc. 3. General Buildings, Offices, Shops, etc. 4. Receiving, Shipping Facilities	859.5 214.9 250.7 376.1	
2a.	SUBTOTAL FOR OTHER DIRECT PLANT INVESTMENT COSTS (PRIMARILY OFFSITE FACILITIES OUTSIDE BATTERY LIM	1,701.2 HITS)	
3.	TOTAL DIRECT PLANT INVESTMENT COST, la + 2a	6,285.4	
4.	INDIRECT FLANT INVESTMENT COSTS 1. Engineering, Overhead, etc. 2. Normal Cont. for Floods, Strikes, etc.	984.9 1,271.4	
4a.	TOTAL INDIRECT FLANT INVESTMENT COST	2,256.3	
5,	TOTAL DIRECT AND INDIRECT PLANT INVESTMENT COST, 3 + 4a	8,541.6	
6.	OVERNLL CONTINGENCY, * of 5	1,708.3	
7.	FIXED CAPITAL INVESTMENT FOR PLANT, 5 + 6	$\times 1.4$	(1975 dollars) inflation (1980 dollars)

TABLE 4.5-11

BETTHATION OF TOTAL PRODUCT GOST FOR SCL PROCESS - CASE >

		\$/¥3 of \$1
1.	Direct Manufacturing Cost (Direct Charges) 1. Raw Materials 2. Direct Operating Labor 3. Utilities 4. Supervision and Clarical 5. Maintenance and Repairs 6. Operating Supplies 7. Laboratory Charge	2.091 0.436 1.083 0.065 1.028 0.208
2.	Indirect Manaufacturing Cost (Fixed Charges) 1. Depreciation 2. Local Taxes 3. Insurance	1.025 0.205 0.103
3.	Plant Overhead	0.608
4.	By-Product Credit	had and a direct department of the second depa
4a.	Total Manufacturing Cost, 1 + 2 + 3 + 4	r.881
5.	General Expenses 1. Administration 2. Distribution and Sales 3. Research and Development	0.413 0.413 0.206
6.	Total Cost of Product, 4a + 5	7.913 (1975 dollars) x 1.4 inflation 11.078 (1980 dollars)

4.5 DCS Process (Dichlorosilane)

The economic analysis activity involves a cost analysis of the DCS process - Case A to produce dichloromilane which is involved in the Hemlock Semiconductor Corporation program form polysilicon.

The cost analysis for the particular technology is based on process design results, such as requirements for raw materials and major process equipment necessary to produce the product from the chemical engineering analysis activity. Primary results issuing from the economic analysis include plant capital investment and product cost which are useful in the analysis of polysilicon production from dichlorosilane.

The cost analysis result for the DCS process (Came A) are presented in Table 4.6-1 including costs for raw materials, labor, utilities and other items composing the product cost (total cost of producing silicon). The tabulation summarizes all of these items to give a total product cost without profit of \$1.39 per kg of DCS (1980 dollars). This product cost without profit includes direct manufacturing cost, indirect manufacturing cost, plant overhead and general expenses.

The product cost represents all cost associated with producing 1 kg of PCS. On top of these costs a predicing company will include some profit. The sales price of the product silicon will actually be the sum of the product cost and a profit for the company. The profit is usually measured in terms of rate of return on the capital investment that the company spent in going into the DCS business. Two profitability methods which are commonly used are the return on original investment (per cent ROI) and discounted cash flow rate of return (per cent DCF).

The cost and profitability analysis summary for this process are presented in Table 4.6-2. The sales price of dichlorosilane at various rates of return for both profitability methods (per cent ROI and DCF) is shown in the lower half of the table. The results indicate a sales price of \$1.47 per kg (1980 dollars) at 15 per cent DCF rate of return on investment.

These cost and profitability results for the DCS process will help the analysis of polysilicon production from dichlorosilane.

The detailed results for the economi nalysis are presented in a tabular format to make it easier to locate cost items of specific interest. The guide for the tabular format is given

below:

	When the Providence of the course of the cou	
٠	Proliminary Moonomic Analysis Activities Table	4.6-8
Ħ	Process Design InputsTable	4.6-4
٠	Base Case Conditions	4.6-5
ě	Raw Material CostTable	4.6-6
٠	Utilitiy Cost	4.6 - 7
٠	Major Process Equipment Cost	4.6-8
•	Production Labor Cost	4.6⊶9
•	Plant InvestmentTable	4.6-10
•	Total Product Cost	4.6-11

The economic analysis provides detailed cost data for raw materials, utilities, labor and major process equipment which are necessary for polysilicon production.

TABLE 4.6-1

BETIMATION OF PRODUCT COST FOR DCS PROCESS

		Cost \$/Kg of DCS (1975 dollars)	Cost \$/Kg of DCs (1980 dollars)
1	Direct Manufacturing Cost (Direct Costs) Raw Materials Direct Operating Labor Utilities Supervision and Clerical Maintenance and Repairs Operating Supplies Laboratory Charge	0.6935	0.9709
2,	Indirect Manufactuing Cost (Fixed Cost) Depreciation Local Taxes Insurance	0.0735	0.1029
з.	Plant Overhead	0.0355	0.0497
4.	General Expenses	0.1205	0.1687
5.	Product Cost Without Profit	0.9230	1.2922

TABLE 4.6-2
COST AND PROFITABILITY ANALYSIS SUMMARY FOR DCS PROCESS

1. 2. 3. 4. 5.	Process			9,780 metric : . dichlerosilan . liquid	•
		Fixed Capit Working Cap (15%)	ital	5.53 Mega 0.83 Mega 6.36 Mega (1975 dollars)	7.74 Maga 1.16 Maga 8.90 Maga (1980 dollars)

E. Return on Original Investment, after taxes (AROI)

		Sales Price \$/Kg of DC8 (1975 dollars)	Sales Price \$/Kg of DCS (1980 dollars)
0%	ROI	. 0.92	1.29
5%	ROI	0.98	1.38
10%	ROI	. 1.04	1.46
15%	ROI	. 1.10	1.85
20%	ROI	. 1.16	1.63
25%	ROI	. 1.22	1.71
30%	ROT	1.28	0.80
40%	ROI	1.40	1.97

7. Discounted Cash Flow Rate of Return, after taxes (DCF)

		\$/Kg of DCS	
		(1975 dollars)	(1980 dollars)
08	DCF	0.92	1.29
5%	DCF,	0.95	1.35
10%	DCF	1.00	1.41
15%	DCF	1.05	1.47
20%	DCF	1.10	1.54
25%	DCF	1.15	1.61
30€	DCF	1.20	1.69
40%	DCF	1.32	1.84

Based on 10 year project life and 10 year straight line depreciation.

TABLE 4.6-3

ECONOMIC ANALYSES: PRELIMINARY ECONOMIC ANALYSIS ACTIVITIES FOR DCS PROCESS

	Prel. Process Economic Activity	Status		Prel. Process Economic Autivity	-ates
1,	Process Design Inputs	*	6.	Production Labor Costs	3
	1. Raw Material Requirements	\$		l. Base Cost Per Man Hour	
	2. Utility Requirements	•		2. Cost/Kg Silicon Per Area	•
	3. Equipment List			3. Total Cost/Kg Silicon	
	4. Labor Requirements				
			7.	Estimation of Plant Investment	
7.	Specify Base Care Jonditions	•		1. Battery Limits Direct Corts	4
	1. Base Year for Costs	•		1. Other Direct Costs	•
	2. Appropriate I. dices for Costs	a		3. Indirect Costs	· e
	3. Addatictal	•		4. Contingency	
				5. Potal Plant Investment	
3.	Raw Material Costs	•		(Fimmā Capital)	
	 Base Cost/Lb of Material 	*			
	Material Cost/Kg of Silicon	•	8.	Est_mation of Total Product Cost	•
	3. Total Cost/Kg of Sillon	9		1. Direct Manufacturing Cost	•
				2. Indirect Manufacturing Cost	•
4.	Thility Costs	•		3. Plant Overhead	\$
	i. Base Cost for Each Utility	4		4. By-Product Credit	•
	3. Utility Cost/%g of Sillcon	#		5. General Expenses	•
	3. Potal Cost/Kg of Silicon	9		6. Total Cost of Product	:
5.	Major Process Equipment Costs	G			
	i. Individual Equipment Cost	ā			
	Cost Index Adjustment	•			
				0 Plan	
				G In Progress	
				* Complete	

TABLE 4.6-4

PROCESS DESIGN INPUTS FOR DCS PROCESS

- 1. Raw Material Requirements
 -Silicon tetrachloride, sinc, lime, argon and nitrogen
 -see table for "Raw Material Cost"
- 2. Utility -electricity, steam. cooling water and process water -see table for "Uti ity Cost"
- 3. Equipment List
 -process vessels, heat exchangers, reactors, etc.
 -see table for "Major Process Equipment Cost"
- 4. Labor Requirements
 -production labor for purification, deposition, electrolysis, etc.
 -see table for "Production Labor Cost"

TABLE 4.6-5 BASE CASE CONDITION FOR DCS PROCESS

1. Capital Equipment

- -January 1975 Cost Index for Capital Equipment Cost
- -January 1975 Cost Index Value = 430

2. Utilities

- -Electrical, Steam, Cooling Water, Nitrogen
- "January 1975 Cost Index (U. S. Dept. Labor)
- -Values determined by literature search and summarized in cost standardization work

3. Raw Material Cost

- -Chemical Marketing Reporter
- -January 1975 Value
- -Raw Material Cost Index for Industrial Chemicals
- -1975 Cost Index Value = 206.9 (Wholesale Price Index, Producer Price Index)

4. Labor Cost

- -Average for Chemical Petroleum, Coal and Allied Industries (1975)
- -skilled \$6.90/hr

5. Update to 1980

- -historically cited 1975 dollars (LSA project)
- -DOE decision to change to 1980 dollars (JPL, 6/22/79)
- -reports to reflect both 1975 and 1980 dollars (JPL, 6/22/79)
- -inflation factor of 1.4 to be used (JPL, 6/22/79)

TABLE 4.6-6

RAW MATERIAL COST FOR DCS PROCESS

	Raw Material	Requirement	\$/1b of material	Cost \$/kg of DCS
1.	M.G. Silicon (Si)	0.348	0.535	0.1862
2.	Silicon Tetrachloride (SiCl ₄ , make-up)	1.987	0.135	0.2682
3.	Liquid Hydrogen	0.048	1.84	0.0883
4.	Copper Catalyst (Cu)	0.005	0.922	0.0046
5.	Hydrate Lime (Ca(OH) ₂)	0.236	0.015 TOTAL	0.0036 0.5509 (1975 dollars) x 1.4 inflation 0.7713 (1980 dollars)

TABLE 4 6-7
UTILITY COST FOR DCS PROCESS

	Utility	Requirement/kg	Cost of Utility	Cost \$/kg of DCS
ı.	Blectricity	0.337 KW HR	0.0324\$/kw hr	0.01092
2.	Steam	12.81 lb	1.35\$/Mlb	0.01730
3.	Cooling water	56.51 gal	0.098/M gal	0.00509
4.	Process water	0.301 gal	0.405\$/M gal	0.00012
5.	Fuel Oil	0.00473 MM Btu	1.40\$/MM Btu	0.00662
			TOTAL	0.04005 (1975 dollars) x 1.4 inflation 0.05607 (1980 dollars)

TABLE 4.6-8

PURCHASED COST OF MAJOR PROCESS EQUIPMENT
FOR DCS PROCESS

	Equipment	Purchased Cost, \$1000
1.	R-01 Hydrochlorination Reactor	81.9
2,	R-02 TCS Redistribution Reactor	14.0
3.	R-03 Wast Neutralizer	15.7
4.	R-04 Waste Combustor	8.0
5.	D-01 Crude TCS Stripping Column	5.40
ß.	D-02 TCS/STC Distillation Column	40.41
7.	D-03 DCS/TCS Distillation Column	94.43
8.	D-04 DCS Distillation Column	41,22
9.	B-01 Silicon Storage Bin with Feed Lock	18.0
10.	T-01 Residue Settling Tank	55.8
11.	T-02 Residue Withdraw Tank	4.0
12.	T-03 Hydrogen Separation Tank	7.9
13.	T-04 Crude TCS Storage Tank	7,5
14,	T-05 TCS Stripper Reflux Drum	0.8
15.	T-06 TCS/STC Distillation Reflux Drum	4.5

TABLE 4.6-8 (centinued) 16. T-07 STC Storage Tank 13.8 6.9 T-08 DC8/TC8 Distillation 17. Reflux Drum 18. T-09 DC8 Distillation 1.2 Roflux Drum 19. T-10 DCS Storage Tank (3) 23.9 ca. 20. T-11 Flue Gam Separation 0.8 Tank 21. T-12 Lime Solution Pre-0.9 paration Tank 22. T-13 Waste Filtrate 0.0 Storage Tank H-OA Crude TCS Condenser 23. 35.7 24. H-02 Ho Gas Preheater 27.2 H-03 STC Vaporizer 25. 54.5 26. H-04 Stripper Condenser 2.0 27. H-05 Stripper Reboiler 3.6 28. H-06 TCS Condenser 53.4 29. H-07 TCS/STC Reboiler 17.7 30. H-08 STC Heat Exchanger 23.4 31. H-09 DCS Condenser 13.4 32. H-10 DCS/TCS Reboiler 14.7 7.2 33. H-11 TCS Cooler 34. H-12 DCS Distillation 1.6 Overhead Condenser 35. H-13 DCS Distillation 0.6 H-14 Wasto Stream Cooler 36. 6.1 37. H 15 STC Superheater 26.7 H-16 H₂ Compressor Intercooler 38. 3.6

TABLE 4.6-8 (continued)

39,	C-01A Hydrogen Feed Compressor, First-stage	26.0	
40.	C-01B Hydrogen Feed Compressor, Second-stage	26.0	
41.	C-02 Hydrogen Circulation Compressor	16.4	
42.	P-01 Feed Tank Blower	8.0	
43.	P-02 Settling Tank Circulation Pump	2.4	
44.	P-04 TCS Reflux Pump	2.7	
45.	P-05 STC Feed Pump	2.7	
46.	P-06 DCS Reflux Pump	1.0	
47.	P-08 DCS Purification Discharge Pump	0.3	
48.	P-09 DCS Pump	6.9	
49.	P-10 Waste Solution Pump	0.5	
50.	P-11 Lime Solution Circulation Pump	0.5	
51.	P-12 Fresh Lime Solution Pump	0.4	
52.	F-01 Silicon Dust Filter	1.6	
53.	F-63 Waste Slurry Filter	5.0	
54.	S-01 Silicon Feed Cyclone	1.4	
55.	E-01 Quench Contact Ejector	1.3	
56.			(1975 dollars) inflation (1980 dollars)

TABLE 4.6-9
PRODUCTION LABOR COST FOR DGS PROCESS

	<u>saction</u>	Labor man-hr/Kg DCS	Labor \$/man-hr	Cost \$/Kg DCS
ı.	Bydrochlorination	0.001394	6.90	0.008929
2.	Purification/Redistribution	0.001941	6.90	0.013393
3.	Waste Treatment	0.000647	6.90	0.004464 0.02679 (1975 dollars) x 1.4 inflation 0.03751 (1980 dollars)

TABLE 4.4-10
ENTIMATION OF PLANT INVESTMENT FOR DCs PRICESS

		Investment \$1000	
1.	DIRECT PLANT INVESTMENT COSTS 1. Major Process Equipment Cost 2. Installation of Major Process Equipment 3. Process Piping, Installed 4. Instrumentation, Installed 5. Flactrical, Installed	891.6 383.4 659.8 169.4 89.2	
la.	6. Progess Buildings, Installed SUBTOTAL FOR DIRECT PLANT INVESTMENT COSTS	89.2 2,282.5	
2.	(PRIMARILY BATTERY LIMIT PACILITIES) OTHER DIRECT PLANT INVESTMENT COSTS 1. Utilities, Installed 2. General Services, Site Development,	428.0 1.07.0	
	Fire Protection, etc. 3. General Buildings, Offices, Shops, atc. 4. Receiving, Shipping Facilities	124.8 187.2	
2a.	SUBTOTAL FOR OTHER DIRECT PLANT INVESTMENT COSTS (PRIMARILY OFFSITE FACILITIES OUTSIDE BATTERY LIMITS)	847.0	
3.	TOTAL DIRECT PLANT INVESTMENT COST, la + 2a	3,129.5	
4.	INDIRECT PLANT INVESTMENT COSTS 1. Engineering, Overhead, etc. 2. Normal Cont. for Floods, Strikes, etc.	490.4 633.0	
4a.	TOTAL INDIRECT PLANT INVESTMENT COST	1,123.4	
5.	TOTAL DIRECT AND INDIRECT PLANT INVESTMENT COST, 3 + 4a	4,252.9	
6.	OVERALL CONTINGENCY, % OF 5	1,275.9	
	FIXED CAPITAL INVESTMENT FOR PLANT, 5 + 6 WORKING CAPITAL INVESTMENT FOR PLANT, % OF 6	$5,528.8$ (19 $\times 1.4$ inf $7,740.3$ (19	lation
	·		

9. TOTAL PLANT INVESTMENT, 7 + 8

TABLE 4.6-11

matination of total product cost for DCs progmss

		8/KG of DCS
1,	Direct Manufacturing Cost (Direct Charges) 1. Raw Materials - from prel. design 2. Direct Operating Labor - from prel. design 3. Utilities - from prel. design 4. Supervision and Clerical 5. Maintenance and Repairs 6. Operating Supplies 7. Laboratory Charge 8. Fatents and Royalties	0.5519 0.0268 0.0400 0.0040 0.0565 0.0113 0.0040
2.	Indiract Manufacturing Cost (Pixed Charges) 1. Depreciation 2. Local Taxes 3. Insurance	0.0568 0.0113 0.0057
3.	Plant Overhead	0.0355
4.	By-Product Credit - from prel. design	
4a.	Total Manufacturing Cost, 1 + 2 + 3 + 4	0.8026
5.	General Expenses 1. Administration 2. Distribution and Sales 3. Research and Development	0.0482 0.0482 <u>0.0241</u>
6.	Total Cost of Product, 4a + 5	0.9230 (1975 dollars) x 1.4 inflation 1.2922 (1980 dollars)

6. SUMMARY - CONCLUSIONS

The following summary-conclusions are made as a result of analyses conducted for new technologies and processes being developed for the production of lower cost milicon for molar cella:

- Analyses of process system properties are important for chemical materials involved in the several processes under consideration for semiconductor and solar cell grade silicon production. Major physical, thermodynamic and transport property data are reported for the following milicon source and processing chemical materials
 - · Silane
 - · Silicon Tetrachloride
 - · Trichlorosilane
 - Dichlorosilane
 - · Silicon Tetrafluoride
 - · Silicon

The property data are reported for critical temperature, critical pressure, critical volume, vapor pressure, heat of vaporization, heat capacity, density, surface tension, viscosity, thermal conductivity, heat of formation and Gibb's free energy of formation. The reported property data are presented as a function of temperature to permit rapid usage in research, development and production engineering.

- 2. Chemical engines analyses involving the preliminary process dealsh of a plant (1000MT/yr capacity) to produce silicon via the techology under consideration were accomplished for the following processes:
 - · UCC Silane Process for Silicon
 - · BCL Process for Silicon Case A
 - · BCL Process for Silicon Case B
 - · Conventional Polysilicon Process (Siemen's Technology)

 - S1I₄ Decomposition Process
 DCS Process (Dichlorosilane)

Major activities in the chemical engineering analyses included base case conditions, reaction chemistry, process flowsheet, material balance, energy balance, property data, equipment design, major equipment list, production labor and lorward for economic analysis. The process design package provided detailed data for raw materials. utilities, major process equipment and production labor requirements necessary for polysilicon production in each process.

- 3. Economic analyses were accomplished for the following processes under consideration for the production of sillcon:
 - · UCC Silane Process for Silicon
 - · BCL Process for Silicon Case A
 - * BCL Process for Silicon Case B
 - Conventional Polysilicon Process (Siemens Technology)
 - * Sil Decomposition Process * DCS Process (Dichlorosilans)

Primary activities in the economic analyses involved procose design inputs, base case conditions, raw material coats utility costs, major process equipment costs and production labor costs in the estimation of plant investment and total product cost.

The cost analysis results for preducing milicon by the UCC milans process (Union Carbide Corporation) are presented including costs for new materials, labor, utilities and other items composing the product cost (total cost of producing silicon) The results indicate a total product cost without profit of \$6.90 (1975 dollars) and \$9.86 (1980 dollars) per kg. For profitability analysis, the results indicate a sales price of \$13 per kg of milicon (1980 dollars) at 15 per cent DCF (discounted cash flow) rate of return on investment.

These cost and profitability results for the UCC silane process indicate that this new technology for producing polysilicon shows good promise for meeting the cost goal of \$14 per kg of silicon waterial (1980 dollars) for solar cells.

The cost analysis results for producing silicon by the BCL process - Case A (Battelle Columbus Laboratories) are presented including costs for raw materials, labor, utilities and other items composing the product cost (total cost of producing silicon). The results indicate a total product cost without profit of \$8.63 (1975 dollars) and \$12.08 (1980 dollars) per kg. The profitability analysis results disclose a sales price of \$13.28 per kg of silicon (1980 dollars) at 5 per cent DCF (discounted cash flow) rate of return on investment after taxes.

These cost and profitability results for the BCL process - Case A indicate that this new technology for producing polysilicon shows promise for meeting the cost goal of \$14 per kg of silicon material (1980 dollars) for solar ceils. In Case A, the process involves two deposition reactors and six electrolysis cells.

The cost analysis resu'ts for producing silicon by the BCL process - Case B (Battelle Columbus Laboratories) are presented including costs for raw materials, labor, utilities and other items composing the product cost

(total cost of producing silicon). The results give a total product cost without profit of \$7.91 (1975 dollars) and \$11.07 (1980 dollars) per kg. Por profitability, the analysis indicates a sales price of \$13.14 per kg of silicon (1980 dollars) at 10 per cent DCF (discounted cash flow) rate of return on investment.

These cost and profitability results for the BCL process - Case B indicate that this new technology for producing polysilicon shows promise for meeting the cost goal of \$14 per kg of silicon material (1980 dollars) for solar cells, In Case B, the process contains one deposition reactor and two electrolysis cells.

7. For the conventional polysilicon process, the cost analysis is based on a poly plant constructed in the 1960's (1965 or earlier) since several existing plants producing semiconductor grade polysilicon in the United States were constructed in the 1960's. The operating costs for the plant are applicable to the time period of interest (such as 1975 and 1980).

The cost analysis results for producing silicon by the conventional Siemens process are presented including costs for raw materials, labor, utilities and other items composing the product cost (total cost of producing silicon). The results (isclose total product cost without profit of \$35.52 - 41.29 (1975 dollars) and \$49.73 - 57.81 (1980 dollars) per kg. The range for product cost reflects low and high electrical costs (1.5-3¢/kw hr for 1975 and 2.1-4.2¢/kw hr for 1980).

The average product cost without profit is estimated at \$38.41 (1975 dollars) and \$53.77 (1980 dollars) per kg for the conventional polysilicon process. This average product costs corresponds to intermediate electrical costs (2.25¢/kw hr for 1975 and 3.15¢/kw hr for 1980). These costs results for the conventional polysilicon process indicate that this Siemens technology using trichlorosilane for producing polysilicon does not show promise for meeting the cost goal of \$14 per kg of silicon material (1980 dollars) for solar cells.

8. The cost analysis results for producing silicon by the Sil, decomposition process are presented including costs for raw materials, labor, utilities and other items composing the product cost (total cost of producing silicon). The results give a total product cost without profit of \$44.64 (1975 dollars) and \$62.50 (1980 dollars) per kg. The profitability results indicate a sales price of \$71.48 per kgof silicon (1930 dollars) at 5 per cent DCF (discounted cash flow) rate of return on investment.

These cost and profitability results for the SiI decomposition process indicate that this new technology for producing polysilicon does not show promise for meeting the cost goal of \$14 per kg of silicon material (1980 dollars) for solar colls.

- 9. Using a hot-wire technique, experimental gas phase thermal conductivity values were determined between 25°C and 350°C for milicon source materials such as milane and halogenated milanes. The accuracy of the values were shown to be 12% by determining values for argon and hydrogen, compounds whose thermal conductivity values have been previously determined.
- 10. Experimental gas phase viscosity values were determined for the halogenated silanes; dichlorosilane, trichlorosilane and tetrafluorosilane. The values were determined in the temperature range 40°C to 200°C using a transpiration method. Nitrogen, a compound for which viscosity values are known, was used to calibrate the apparatus. The calibration studies showed that the values obtained were accurate to 12% throughout the temperature range.
- 11. Studies were conducted to develop an efficient method for the generation of SiF, from hexafluorosilicic acid, a readily available by-product of the phosphate fertilizer industry. This included investigation of such parameters as conditions for precipitation of SiF, procursors (Na₂SiF and BaSiF₆), temperature for thermal decomposition of the salts, heating time required and optimum flow rates. Precipitation of the salts, Na₂SiF₆ or BaSiF₆, with NaCl, NaF, BaCl₂, or BaF₂ followed by thermal decomposition at temperatures above 500°C proved to be an efficient method for the generation of SiF₄.

A1. ADDITIONAL CHIMICAL INGINIBRING ANALYSIS

A1.1 Silane Process - Case A

The chemical engineering analysis activity of Silane Process - Case A (Regular Process Storage) involves a preliminary process design of a plant to product silane for silloon.

The Silane Process-Case A involves several processing operations of hydrogenation, distillation, redistribution reaction, stripping and absorption. The process flowsheet is shown in Figure Al.1-1. This flowsheet was received from Union Carbide.

A summation of the salient feature of Case A is shown below:

CARE A

Process	
Process Flowsheet	
Process Chemistry and Equilibrium	From Union Carbide
Intermediate Product Storage Considerations Major Process Equipment	

The detailed status sheet is shown in Table Al.1-1, and is representative of the various subitems that make up the preliminary design activity. The results from the preliminary process design are presented in a tabular format similar to previous design results for alternate processes to produce silicon. Note that in this process results are per pound of silane versus other processes represented as per kilogram of silicon. The silane plant size assumes a 90% conversion of silane to silicon.

The guide to the tables for Case A is given below:

Base Case Conditions	A1.1-2
Reaction Chemistry	A1.1-3
Raw Material Requirement Table	A1.1-4
Utility Requirements	A1.1-5
Major Process Equipment	
Production Labor Requirements	A1.1-7

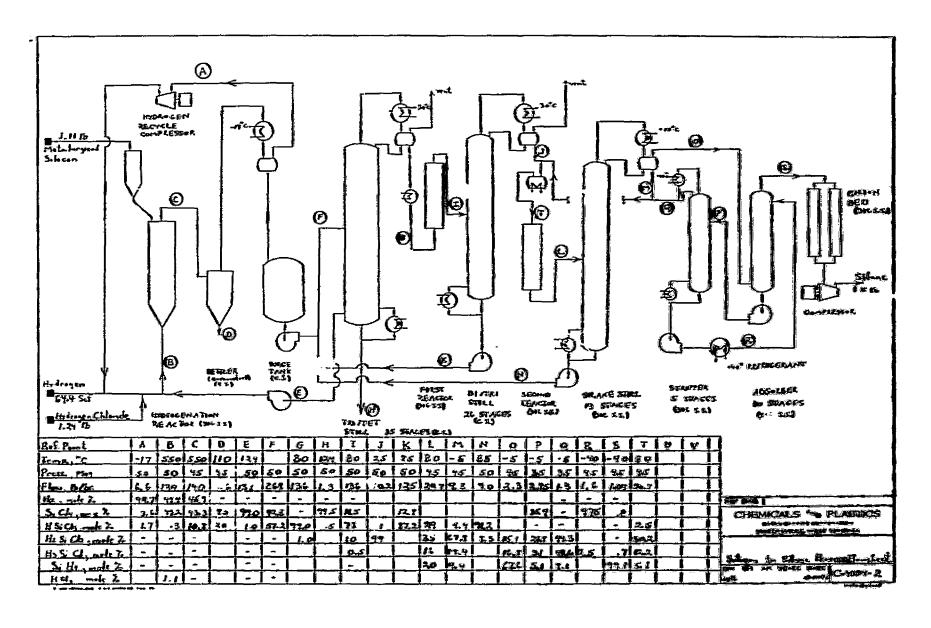


Figure Al.1-1 Process Flow Sheet for Silane Process-Case A (Provided by Union Carbide)

CASB A

TABLE AL.1-1 CHEMICAL ENGINEERING ANALYSES: PRELIMINARY PROCESS DESIGN ACTIVITIES FOR SILANE PROCESS - CASE A

		Prel. Process Design Activity	Status		Prel. Process Resign Autivity	States
	1.	Specify Base Case Conditions 1. Plant Size 2. Product Specifics 3. Additional Conditions	•	7.	Equipment Design Calculations 1. Storage Vessels 2. Smit Operations Equipment 3. Process Data (P, T, rate, etc.) 4. Additional	6 3 0
	2.	Define Reaction Chemistry 1. Reactants, Products 2. Equilibrium	•	8.	List of Major Process Bunipment 1. Size 2. Type	9 0 0
64 60 20	3.	Frocess Plow Diagram 1. Plow Sequence, Unit Operations 2. Process Conditions (T, P, etc.) 3. Environmental 4. Company Interaction (Technology Exchange)		8a.	3. Materials of Construction Major Technical Pactons (Potential Problem Areas) 1. Materials Compatibility 2. Process Conditions Limitations 3. Additional	•
	4.	Material Balance Calculations 1. Paw Materials 2. Products 3. By-Products		9.	Production Labor Requirements 1. Process Technology 2. Production Wolume	0
	5.	Energy Balance Calculations 1. Heating 2. Cooling 3. Additional		10.	Porwand for Boomonic Analysis	•
	6.	Property Data 1. Physical 2. Thermodynamic 3. Additional			0 Plan © In Progness • Complete	

7

CASH A

TABLE Al.1-2

Base care conditions for silane process - Case A

- 1. Plant Size; Allow for 10% lesses of silane in production of silicon
 - 1270 metric tons/year of silane
 - Solar cell grade silicon
- 2. Hydrogenation Reaction
 - Metallurgical grade silicon, hydrogen, to produce trichlorosilane (FCS) make-up hydrogen chloride used and recycle silicon tetrachloride (TET)
 - Copper catalysed
 - Fluidized bed
 - 650°C, 50 PSIG
 - 15.8% conversion of SiCl4 (Union Carbide flowshest)
- TCS Redistribution Reaction
 - TCS from hydrogenation produces dichlorosilane (DCS)
 - Catalytic redistribution of TCS with tertiary amine ion exchange resin.
 - Liquid phase 50 PSIG, 80°C.
 - Conversion a function of inlet concentration per Figure IIA-2 (Union Carbide equilibrium)
 - Conversion from pure TCS feed is about 10% to DCS (example)
- DCS Redistribution Reaction

 - DCS produces SiH4 (silans)
 Catalytic redistribution of DCS with tertiary amine ion exchange resin.
 - Gas phase 60-80°C
 - Conversion a function of inlet concentration per Figure IIA-1.1 (Union Carbide equilibrium)
 - Conversion from pure DCS feed is about 14% to Silane (example)
- 5. Recycles
 - Unreacted chlorosilanes separated by distillation and recycled
- Silane Purification
 - Chlorosilanes removed by absorption in -40°C SiCl, (Tet)
 - Trace contaminants removed by carbon adsorption
- 7. Operating Ratio
 - Approximately 90% utilization
 - Approximately 7880 hour/year production
- 8. Storage Considerations
 - Feed materials (two week supply)
 - Product (two week supply)
 - Process (several days)

CASH A

TABLE AL.1-8

REACTION CHEMISTRY FOR SILAND PROCESS - CASE A

- 1. Hydrogenation Reaction
 - 3 81Cl₄ + 81 + 2H₂ * 481HCl₃
- 2. Trichlorosilane Redistribution Reaction
 - 2 Sincl₃ * Sin₂cl₂ + Sicl₄
- 3. Dichlorosilane Redistribution Reaction
 - 3 Sim₂Cl₂ Distillation 2SimCl₃ + Sim₄

Note

- 1. Reaction 1 Product contains H₂, SiCl₄, SiHCl₃, SiH₂Cl₂ (trace), other trace colorides
- 2. Reaction 2 Product contains SiHCl3, SiCl4, SiH2Cl2, SiH3Cl
- 3. Reaction 3 Product contains SiH_2Cl_2 , $SiHCl_3$, $SiCl_4$, SiH_3Cl , SiH_4

CAST A

TABLE AL.1-4

naw maderial requirements for Silane Process - Cash A

	Row Material	Requirement <u>16/16 of Silane</u>
1.	Anhydrons HCl	1.239
2.	Rydrogen	.362
3.	Caustic (50%)	2.448
4.	M.G. Silionn	1.11

SMI A

TABLE Al.1-5

UTILITY REQUIREMENTS FOR SILANS PROCESS - CASE A

<u>U</u>	tility/Punction	Requirements/1b of	#11ane	Product
1.	Electricity 1. All pump and compressor motors (24)	(.253)	. 253	KW-HR
2.	Stram 250 Psia 1. #1 Pistillation Column Preheater 2. #1 Distillation Column Reboiler 3. #2 Pistillation Column Reboiler 4. #2 Redistribution Reactor Preheater 5. #3 Distillation Column Preheater 6. #3 Distillation Column Reboiler 7. #4 Distillation Column Reboiler 8. Wasta Treatment	(6.96) (81.18) (91.77) (3.0) (3.62) (3.29) (.41) (.11)	190.34	lbs
3.	Cooling hater (10-120°F) 1. #1 Fistillation Column Condenser 2. #2 Distillation Column Condenser	(145.12) (22.09)	168.12	gallons
4.	Process Water (90°F) 1. Waste Treatment	(8.22)	8.22	gallons
5.	Refrigerant (23°F) 1. #4 Distillation Column Feed Tank	(27.1)	27.1	BTO
6.	Refrigerant (5°F) 1. #3 Distillation Column Overhead Receiver	(79.1)	79.1	BTU
7.	Refrigerant (-7°F) 1. #4 Distillation Column Overhead Receiver	(26.4)	26.4	BTO
8.	Refrigerant (-20°F) 1. #3 Distillation Column Condenser 2. #4 Distillation Column Condenser	(2058.0) (245.2)	2303.2	BTU

CABE A

9.	Refrigerant (-30°F)		30788.0 BTU
	1. TCS Reactor Recycle Gas Condenser	(30788.0)	
10.	Refrigerant (-40°P)		280.6 BTU
	1. #2 Redistribution Reactor Con-		
	dengate Receiver	(192.2)	
	2. Silane Product Storage	(88.4)	
11.	Refrigerant (~50°F)		3503.2 BTU
	1. #2 Radistribution Reactor Gas Condensor	(2986.0)	
	2. Froduct Silane Condenser	(137.9)	
	3. Absorbent Cooler	(379.3)	
12.	High Temperature Heat Exchange Pluid		3.324 x 10 ⁴ BTU
	1. TCS Reactor Recycle Gas Hauter	(6.591×10^3)	
	2. HCl Vaporizer	(4.466×10^2)	
	3. Tet Vaporizer	(2.464×10^4)	
	4. Hell; Nitrogen to Regnerate Whar. Adsorbers	(70.95)	
	5. TCS Reactor	(1.491×10^3)	
13.	Nitrogen		5.54 SCF
	1. Regenerate Charcoal Adsorbers	(5.54)	

TABLE AL.1-E

LIST OF MAJOR PROCESS BOSTPMENT FOR SILANE PROCESS - CASE A

						Matterfials
		Type	Function	Duty	Size	of Construction
1.	(T1)	M.G. Silion Storage Hopper	Raw Material Storage	2 weeks storage	1.363 x 10 ⁴ gallon	s CS
2.	(T2)	Hydrogen Storage Tank	Raw Material Storage	8 hours backup for pipeline failure	9.161 x 10 ⁴ gallos 250 PSTA (spherios	
3.	(T 3)	Liquid ECL Storage TAnk	Raw Material Storage	2 weeks storage	1.612 × 10 ⁴ gellom 250 PSIA -50 ⁶ F (spherical)	s Michel Steel
4.	(T4)	Recycle TRT Stonage	For TCS Beactor Feed	2 days storage	1.985 x 10 ⁵ gallow 65 PSIA	e CS
5.	(T5)	TCS Reactor Off- Gas Plash Tank	Phase Separation		1 ft. diameter by a long, 65 PSTA, 6°P 65 PSTA	ift. es
6.	(T6)	TCS/TET Storage	Peed Distillaton Column #1	2 Days bold-up	1.966 x 10 ⁵ gallons 65 PESA	s CS
7.	(T7)	#1 Distillation Column Condensate Aconmulator	Reflex feed; column Control	20 minutes hold-up	4.88 x 10 ³ gallons 65 ISHA	C9
8.	/T8)	#1 Redistribution Reactor Feed Tank	Hold—up and feed Reactor	2 days hold-up	2.266 x 10 ⁵ gallon 65 PSUA	s (cs
9.	(T9)	#1 Redistribution Reactor Product Tank	Hold-up and feed #2 Distillation Column	2 days hold-up	2.21 x 10 ⁵ gallons 65 PSTA	©\$

	10.	(T10)	#2 Distillation Column Condensate Accumulator	Reflux feed; column Control	20 minutes hold-up	745 gallons 65 PSIA	SS
	11.	(T11)	#2 Redistribution Reactor Feed Tank	Hold-up and feed Reactor	2 days hold-up	1.891 x 10 ⁴ gallons 65 PSIA	SS
	12.	(T12)	#2 Redistribution Reactor Product Tank	Hold-up and feed #3 Distillation Column	2 days hold-up 6.8 x 10 ⁴ BTU/hr	3.46 x 10 ⁴ gallons -40°F, 60 PSIA	SS
	13.	(T13)	#3 Distillation Column Condensate Accumulator	Reflux feed; phase Separation; column control	20 minutes hold-up	194 gallons 5°F, 60 PSIA	SS
366	14.	(T14)	#3 Distillation Column Condensate Tank	Hold-up and recycle feed to #2 Redistribu- tion Reactor	2 days hold-up 2.81 x 10 ⁴ BTU/hr	1.7 x 10 ⁴ gallons 60 PSIA, 5°F	SS
	15.	(T15)	#4 Distillation Column Feed Tank	Surge between absorber and distillation	2 days hold-up 9.63 x 10 ³ BTU/hr	4.69 x 10 ³ gallons 60 PSIA	SS
	16.	(T16)	#4 Distillation Column Condensate Accumulator	Reflux feed; column control	20 minutes hold-up	16 gallons 50 PSIA,-7°F	SS
	17.	(T17)	#4 Distillation Column Condensate Tank	Hold-up and recycle to #2 Redistribution Reactor	2 days hold-up 9.4 x 10 ³ BTU/hr	2.55 x 10 ³ gallons 50 PSIA, -7°F	SS
	18.	(T18)	Waste Tank	Collect waste for Treatment and disposal	2 week storage	1.378 x 10 ⁴ gallons 65 PSIA	cs
	19.	(T19)	Absorber Feed Tank	Feed TET to absorber	2 days storage	2.44 x 10 ³ gallons 50 PSIA	SS
	20.	(T20)	Silane Storage	Final Product storage	1 week storage 3.14 x 10 ⁴ BTU/hr	1.522 x 10 ⁴ gallons -40°F, 250 PSIA	SS

21.	(T21)	Caustic Storace	Raw Material Storage	2 weeks storage	2.304×10^4 gallons	S S
22.	(H1)	TCS Reactor Recycle Gas Heater	Heat Recycle gas and Hydrogen to 550°C	2.342 x 10 ⁶ BTU/hr	752 ft ² 65 PSIA	cs
23.	(H2)	HCl Vaporizer	Heat Reactant to 550°C	1.587 x 10 ⁵ BTU/hr	34 ft ² 65 PSIA	CS
24.	(H3)	TET Va, orizer	Heat Reactant to 550°C	8.755 x 10 ⁶ ETU/nr	2381 ft ² 65 PSIA	CS
25.	(H4)	TCS Reactor Re- cycle Condenser	Phase separation; Recycle hydrogen	1.094 x 10 ⁷ BTU/hr	1882 ft ² 65 PSIA	CS/SS
26. 4.	(H5)	#1 Distillation Column Preheater	Preheat distillation feed to bubble point	2.044 x 10 ⁶ BTU/hr	164 ft ² 250 PSIA	cs
ô ₂₇ .	(H6)	#1 Distillation Column Condenser	Provide Reflux to Column	1.296 x 10 ⁷ BTU/hr	3189 ft ² 65 PSIA	cs
28.	(H7)	#1 Distillation Column Reboiler	Provide vapor to Column	2.382 x 10 ⁷ BTU/hr	2818 ft ² 250 PSIA	cs
29.	(H8)	#2 Distillation Column Condenser	Provide Reflux to column	1.96 x 10 ⁶ BTU/hr	956 ft ² 65 PSIA	CS/SS
30.	(H9)	#2 Distillation Column Reboiler	Provide Vapor to Column	2.693 x 10 ⁷ B fU /hr	2514 ft ² 250 PSIA	cs
31.	(H10)	#2 Redistribution Reactor Feed Vaporizer	Vaporize Reactants for Reactor	8.81 x 10 ⁵ BTU/hr	78 ft ² 250 PSIA	cs/ss
32.	(H11)	#2 Redistribution Reactor Product Condenser	Condense Vapor for hold-up storage	1.06 x 10 ⁶ BTU/hr	306 ft ² 60 PSIA	CS/SS
33.	(H12)	#3 Distillation Column Preheater	Vaporize and preheat feed to column	1.06 x 10 ⁶ BTU/hr	66 ft ² 250 PSIA	CS/SS

34.	(H13)	#3 Distillation Column Condenser	Provide Column Reflux (Partial Condenser)	7.312 x 10 ⁵ BTU/hr	593 ft ² 60 PSIA	CS/SS
35.	(H14)	#3 Distillation Column Reboiler	Provide Vapor to Column	9.64 x 10 ⁵ BFU/hr	84 ft ² 250 PSIA	CS/SS
36.	(H15)	Silane Condenser	Condenser Final Pro- duct for storage	4.9 x 10 ⁴ BTU/hr	53 ft ² 250 PSIA	CS/SS
37.	(H16)	#4 Distillation Column Condenser	Provide Reflux	8.71 x 10 ⁴ BTU/hr	84 ft ² 50 PSIA	cs/ss
38.	(H17)	#4 Distillation Column Reboiler	Provide Vapor to Column	1.2 x 10 ⁵ BTU/hr	13 ft ² 250 PSIA	CS/SS
Ĉ 39.	(H18) Absorber Pre- cooler	Cool TET for absorp- tion column	1.35 x 10 ⁵ BTU/hr	35 ft ² 60 PSIA	CS/SS	
40.	(H19)	Nitrogen Heater	Heat Nitrogen to re- generate Charcoal Adsorbers	2.52 x 10 ⁴ BTU/hr	14.1 ft ²	cs
41.	(Pl)	TCS Reactor Off Gas Recycle Com- pressor	Circulate Recycle Gas to Reactor	1.36 x 10 ³ SCFM	26.5 Horsepower 75 PSIA Discharge	CS*
42.	(P2)	#1 Distillation Column Feed Pump	Feed Column	136.5 gpm	106 PSI; 14.5 BHP	CS*
43.	(P3)	#1 Distillation Column Overheads Pump	Provide Reflux and remove overhead product	244 дрш	92.3 PSI; 22.5 PHP	CS*
44.	(P4)	#1 Distillation Column Bottoms Pump	Remove Bottoms Product to TET storage tank	69 gpm	106 PSI; 7.3 BHP	CS*

	45.	(P5)	Process Water Feed Pump	Feed Process Water to Waste Treatment	48.6 gpm	82.5 PSI; 4 BHP	CS*
	46.	(P6)	Caustic Feed Pump	Feed Raw Material to waste treatment	1 gpm	118 PSI; ½ BHP	SS
4	47.	(P7)	#1 Redistribu- tion Reactor Feed Pump	Feed TCS to Reactor	79 gpm.	106 PSI; 8.4 BHP	SS
	48.	(P8)	#2 Distillation Column Feed Pump	Feed TCS/DCS still	76.6 gpm	92.3 PSI; 7.1 BHP	SS
402	49.	(P9)	#2 Distillation Column Overheads Pump	Provide Reflux and Remove Overhead Product	37.3 gpm	92.3 PSI; 3.4 BHP	SS
-	50.	(P10)	#2 Distillation Column Bottoms Pump	Remove Bottoms Product to TCS/TET storage tank	66.7 gpa	106.3 PSI; 7.1 BPH	SS
	51.	(P11)	#2 Redistribution Reactor Feed Pump	Feed DCS to Reactor	13.4 gpm	130 PSI; 1.7 BHP	SS
	52.	(P12)	#3 Distillation Column Feed Pump	Feed Silane Still	12 gpm	87.3 PSI; 1 BHP	SS
	53.	· · · · · · · · · · · · · · · · · · ·	Provide Reflux; Remove Overhead Product	9.7 дря	87.3 PSI; 1 BHP	SS	
54. 55.	54.	(P14)	#3 Distillation Column Bottoms Pump	Remove Bottoms Pro- duct to TCS/TET Tank	5.2 дры	106.3 PSI, ½ BHP	SS
	55.	(P15)	#4 Distillation Feed Pump	Feed TET Stripper	1.6 gpm	77.3 PSI, 1/4 BEP	SS

^{*} Includes incremental higher cost for special purity requirements.

	56.	(P16)	#4 Distillation Column Overhead Pump	Provide Reflux, Remove Overhead Product	1 дря	77.3 PSI; ¼ BHP	SS
	57.	(P17)	#4 Distillation Column Bottoms Pump	Remove Bottoms Product to Absorber Feed Tank	l gpm	91.3 PSI; ½ BEP	SS
	58.	(P18)	#4 Distillation Condensate Re- cycle Pump	Recycle Condensate back to #2 Redistri- bution Reactor	l gpm	106.3 PSI; ½ BHP	SS
	59.	(P19)	Silane Product Compressor	Liquefy Silane for Storage	66 SCPM	250 PSIA Discharge 6.5 HP	SS
403	60.	(P20)	Waste Feed Pump	Distillation Wastes to Waste Treatment	1 gpm	76.3 PSI; ABHP	cs
	61.	(P21)	TCS Reactor Feed Pump	Feed TET to Reactor	69 дря	92.3 PSI; 6.4 BEP	CS*
	62.	(P22)	#3 Distillation Condensate Re- cycle Pump	Recycle Condensate back to #2 Redistribu- tion reactor	5.9 gpa	92.3 PSI; ½ BHP	5 5
	63.	(P23)	Waste Collection Pump	Distillation Wastes to Waste Tank	1 gpa	87.3 PSI; ¼ BEP	cs
	64.	(P24)	Absorber Feed Pump	Feed Cold TET to Absorption Column	1 gpm	87.3 PSI; ½ BHP	SS
	65.	(C1)	#1 Distillation Column	Separate TET from	95,220 lb/hr of feed	7.56 ft. diameter 100 ft. tall, 50 trays	CS

TABLE Al.1-6 (continued)

66.	(C2)	#2 Distillation Column	Separate TCS from	48, 321 lb/hr of feed	10.6 ft. Diameter 136 ft. tall, 68 trays	cs
67.	(C3)	#3 Distillation Column	Separate Silane from other Chlorosilanes	7344 lb/hr of feed	2.01 ft. Diameter 29 ft. tall, 29 trays	SS
68.	(C4)	#4 Distillation Column	Strip TET for use in absorber	1007.7 lb/hr of feed	1.04 ft. Diameter 28.5 ft. tall, 38 trays	SS
69.	(C5)	Silane Absorber	Absorb Chlorosilane from Silane	819.3 lb/hr of vapor feed	0.823 ft. Diameter 12 ft. tall, 16 trays	SS
70.	(C6)	Charcoal Adsorber	Activated Carbon Adsor- btion of Silane to re- move Trace Chlorosilane	366 lb/hr of vapor feed	1 ft. Diameter 7 ft. tall (2), 623 lbs of carbon	SS
71.	(Rl)	TCS Fluidized Bed Reactor	Produces TCS from TET M.G.Silicon, and H ₂		6.26 ft. in diameter 26.5 ft. tall, 481 tubes 1", 16' long	SS
72.	(R2)	#1 Redistribu- tion Reactor	Redistribute TCS to DCS		2' Diameter by 15 ft. tall 1042 lbs catalyst	SS
73.	(R3)	#2 Redistribu- tion Reactor	Redistribute DCS to Silane		2.34' Diameter by 35 ft.tall 1667.2 lbs catalyst	SS
74.	(Al)	Fines Separator	Remove Silicon Fines carried over with TCS Reactor Off-gas		Standard design 30° Diameter	SS
75.	(A2)	Waste Treatment	Discharge innocuous effluent		l column for absorption + l heat exchanger to vapor- ize feed	56
76.	(A3)	Hydrogen Flare	Dispose of Hydrogen from Waste Treatment		30 ft. stack 6° Diameter	CS
	67. 68. 69. 70. 71. 73.	67. (C3) 68. (C4) 69. (C5) 70. (C6) 71. (R1) 72. (R2) 73. (R3) 74. (A1) 75. (A2)	Column 67. (C3) #3 Distillation Column 68. (C4) #4 Distillation Column 69. (C5) Silane Absorber 70. (C6) Charcoal Adsorber 71. (R1) TCS Fluidized Bed Reactor 72. (R2) #1 Redistribution Reactor 73. (R3) #2 Redistribution Reactor 74. (A1) Fines Separator 75. (A2) Waste Treatment	Column Column Column Separate Silane from other Chlorosilanes 68. (C4) #4 Distillation Strip TET for use in absorber 69. (C5) Silane Absorber Absorb Chlorosilane from Silane 70. (C6) Charcoal Adsorber Activated Carbon Adsorbtion of Silane to remove Trace Chlorosilane 71. (R1) TCS Fluidized Produces TCS from TET M.G.Silicon, and H2 72. (R2) #1 Redistributor Redistribute TCS to DCS 73. (R3) #2 Redistributor Redistribute DCS to Silane 74. (A1) Fines Separator Remove Silicon Fines carried over with TCS Reactor Off-gas 75. (A2) Waste Treatment Discharge innocuous effluent	Column Column	Column DCS of feed 136 ft. tall, 68 trays 67. (C3) #3 Distillation Separate Silane from other Chlorosilanes of feed 29 ft. tall, 29 trays 68. (C4) #4 Distillation Strip TET for use in absorber in absorber of feed 28.5 ft. tall, 35 trays 69. (C5) Silane Absorber Absorb Chlorosilane from Silane of vapor feed 12 ft. tall, 16 trays 70. (C6) Charcoal Adsorber Activated Carbon Adsorbetion of Silane to remove Trace Chlorosilane Dead Reactor RET M.G.Silicon, and H2 71. (R1) TCS Fluidized Produces TCS from TET M.G.Silicon, and H2 72. (R2) #1 Redistribu- Redistribute TCS to DCS 73. (R3) #2 Redistribu- Redistribute DCS to Silane Carbon Picton Reactor Silane 74. (A1) Fines Separator Remove Silicon Fines carried over with TCS Reactor Off-gas 75. (A2) Waste Treatment Discharge innocuous effluent Dispose of Bydrogen 30 ft. stack

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CASE A TABLE Al.1-7

PRODUCTION LABOR REQUIREMENTS FOR SILANE PROCESS-CASE A

	Unit Operation	Туре	Skilled Labo Per Day Po	or,Man Hours or lb.Silane	Semiski Per Day	lled Labor Per lb.silane
1.	TCS Production	В	65	.0085		
2.	Hydrogen Recycle	Ç	18	.0023		
з.	Raw Material Vaporization	C	50	.0065		
4.	TCS Condensation	C	50	.0065		
5.	TCS/TET Separation	С	62	.0081		
6.	#1 Redistribution Reactor	С	49	.0064		
7.	DCS/TCS Separation	С	52	.0068		
8.	#2 Redistibuiton Reactor	С	32	.0042		
9.	Silane Distillation	C	32	.0042		
10.	Silane Absorption	C	28	.0036		
11.	Silane Purification (adsorption)	A	36	.0047		
12.	Silane compression	В	23	.003		
13.	Silane Condensation	В	23	.003		
14.	Materials Handling	λ			48	.0063
15.	Waste Treatment	В	60	.0078		
16.	Silicon Fines Separation	A	15	.002		
	TOTAL		 - 595	.0776	48	.0063

NOTES:

^{1.} A Batch Process of Multiple Small Units

B Average Process

c Automated Process

^{2.} Man hours/day Unit from Figure 4-6, Peters and Timmerhaus (7).

A1.2 Silane Process - Case B

The chemical engineering analysis of the Silane Process - Case B (Minimum Process Storage) involves a preliminary process design of a plant to produce silane for silicon.

The Silane Process - Case B involves several processing operations of hydrogenation distillation, redistribution, distillation, redistribution reaction, stripping and absorption. The flowsheet received from Union Carbide, upon which the design is based, is shown in Figure A1.2-1.

A summation of the important features of CASE B is presented in the following table:

CASE B

Process	Silane (Union Carbide)
Plant Size	.1270 MT/year of Silane
Process Flow Sheet	Original Received from
	Union Carbide
Process Chemistry & Equilibrium	From Union Carbide
Intermediate Product Storage Considerations	.Minimum
Major Process Equipment	.58 pieces of Process
	Equipment

The results from the preliminary process design (CASE B) are summarized in a tabular format parallel to those representing Case A. These tables are represented by the following guide to enable the reader to quickly locate items of interest.

Base Case Conditions	.Table	A1.2-2
Reaction Chemistry	.Table	A1.2-3
Raw Material Requirement		
Utility Requirements		
Major Process Equipment	.Table	A1.2-6
Production Labor Requirements		

Figure Al.2-1 Process Plow Sheet for Silane Process -CASE B (Provided by Union Carbide)

TABLE Al.2-1 CHEMICAL ENGINEERING ANALYSES: PRELIMINARY PROCESS DESIGN ACTIVITIES FOR SILAME PROCESS -CASE B

	Prel. Process Design Activity	<u>Status</u>	Prel. Process Design Activity Status
1.	Specify Base Case Conditions	•	7. Equipment Design Calculations
	l. Plant Size	•	1. Storage Vessels
	2. Product Specifics	•	2. Unit Operations Equipment
	3. Additional Conditions	•	<pre>3. Process Data (P, T, rate, etc.)</pre>
			4. Additional •
2.	Define Reaction Chemistry	•	
	1. Reactants, Products	•	8. List of Major Process Equipment
	2. Equilibrium	•	1. Size
	-		2. Type ●
3.	Process Flow Diagram	•	3. Materials of Construction
	1. Flow Sequence, Unit Operations	•	
	2. Process Conditions (T, P, etc.)	•	8a. Major Technical Pactors
	3. Environmental	•	(Potential Problem Areas)
	4. Company Interaction	•	l. Materials Compatibility
	(Technology Exchange)		2. Process Conditions Limitations
			3. Additional
4.	Material Balance Calculations	•	
	l. Raw Materials	•	9. Production Labor Requirements
	2. Products	•	1. Process Technology
	3. By-Products	•	2. Production Volume
5.	Energy Balance Calculations	•	10. Forward for Economic Analysis
	1. Heating	•	
	2. Cooling	•	
	3. Additional	•	
6.	Property Data	•	0 Plan
	1. Physical	•	€ In Progress
	2. Thermodynamic	•	● Complete
	3. Additional	•	

TABLE Al.2-2

BASE CASE CONDITIONS FOR SILANN PROCESS-CASE B

- 1. Plant Size
 - Allow for 10% losses of silane in production of silicon
 - 1270 metric tons/year of silane
 - Solar cell grade silicon
- 2. Hydrogenation Reaction
 - Metallurgical grade silicon, hydrogen, to produce trichlorosilane (TCS) make-up hydrogen chloride used and recycle silicon tetrachloride (TET)
 - Copper catalyzed
 - Fluidized bed
 - 550°C, 50 PSIG
 - 15.8% conversion of SiCl4 (Union Carbide flowsheet)
- 3. TCS Redistribution Reaction
 - TCS from hydrogenation produces dichlorosilane (DCS)
 - Catalytic redistribution of TCS with tertiary amine ion exchange resin.
 - Liquid phase 50 PSIG, 80°C.
 - Conversion a function of inlet concentration per Figure IIA-2 (Union Carbide equilibrium)
 - Conversion from pure TCS feed is about 10% to DCS (example)
- 4. DCS Redistribution Reaction

 - DCS produces SiH_A (silane) Catalytic redistribution of DCS with tertiary amine ion exchange resin.
 - Gas phase 60-80°C
 - Conversion a function of inlet concentration per Figure IIA-1.1 (Union Carbide equilibrium)
 - Conversion from pure DCS feed is about 14% to Silane (example)
- 5. Recycles
 - Unreacted chlorosilanes separated by distillation and recycled
- 6. Silane Purification
 - Chlorosilanes removed by absorption in -40°C SiCl_A(Tet)
 - Trace contaminants removed by carbon adsorption
- 7. Operating Ratio
 - Approximately 90% utilization
 - Approximately 7880 hour/year production
- 8. Storage Considerations
 - Feed materials (two week supply)
 - Product (two week supply)
 - Process (several days)

TABLE Al.2-3

REACTION CHEMISTRY FOR SILANS PROCESS - CASE B

- 1. Hydrogenation Reaction
 - $3 \operatorname{sicl}_4 + \operatorname{si} + 2\operatorname{H}_2 \stackrel{*}{\leftarrow} 4\operatorname{sincl}_3$
- 2. Trichlorosilane Redistribution Reaction
 - $2 \operatorname{siHCl}_3 \stackrel{*}{\Rightarrow} \operatorname{siH}_2 \operatorname{Cl}_2 + \operatorname{siCl}_4$
- 3. Dichlorosilane Redistribution Reaction
 - 3 simpola Distillation 2siHCl3 + SiH4

Note

- Reaction 1 Product contains H₂, SiCl₄, SiHCl₃, SiH₂Cl₂ (trace), other trace chlorides
- 2. Reaction 2 Product contains $SiHCl_3$, $SiCl_4$, SiH_2Cl_2 , SiH_3Cl
- 3. Reaction 3 Product contains SiH2Cl2, SiHCl3, SiCl4, SiH3Cl, SiH4

CASE B

TABLE A1.2-4

RAW MATERIAL REQUIREMENTS FOR SILANE PROCESS-CASE B

Raw Material		Requirement lb/lb of Siland		
ı.	Anhydrous HCl	1.239		
2.	llydrogen	.362		
3.	Caustic (50%)	2.448		
4.	M.G. Silicon	1.11		

TABLE A1.2-5
UTILITY REQUIREMENTS FOR SILANE PROCESS - CASE B

	Utility/Function	Requirements/1b of	Silane Product
1.	Blactricity 1. All pump and Compressor		.212 KW-HR
	Motors (16)	.212)	
2.	Steam 250 Psia		186.72 lbs
	1. #1 Distillation Column Preheater	(6.96)	
	2. #1 Distillation Column Reboiler	(81.18)	
	3. #2 Distillation Column Reboiler	(91.77)	
	4. #2 Redistribution Reacotr Preheater	(3.0)	
	5. #3 Distillation Column Reboiler	(3,29)	
	6. #4 Distillation Column Reboiler	(0.41)	
	7. Waste Treatment	(0.11)	
3.	Cooling Water (10-120°F)		168.12 gallons
	1. #1 Distillation Column Condenser	(146.12)	
	2. #2 Distillation Column Condenser	(22.09)	
4.	Process Water (90°F)		8.22 gallons
	1. Waste Treatment	(8.22)	-
5.	Refrigerant (-20°F)		2303.2 BTU
	1. #3 Distillation Column Condenser	(2058.0)	
	2. #4 Distillation Column Condenser	(245.2)	
6.	Refrigerant (-30°F)		30788.0 BTU
	1. TCS Reactor Recycle Gas Condenser	(30788.0)	
7.	Refrigerant (-40°F)		25.26 BTU
	1. Silane Product Storage	(25.26)	
8.	Refrigerant (-50°F)		517.2 BTU
	1. Product Silane Condenser	(137.9)	
	2. Absorbent Cooler	(379.3)	
9.	High Temperature Heat Exchange Fluid	2	3.324 x 10 ⁴ BTU
	l. TCS Reactor Recycle Gas Heater	(6.591×10^3)	
	2. HCl Vaporizer	(4.46 x 10 ²)	
	3. Tet Vaporizer	(2.464 x 10 ⁴)	
	4. Heat Nitrogen to Regenerate Char.		
	Adsorber s	(70.95)	
	5. TCS Reactor	(1.491×10^3)	

TABLE Al.2-5 (Continued)

Utility/Function Requirements/1b of Silane Product 10. Nitrogen 1. Regenerate Chargoal Adsorbers (5.54)

TABLE Al.2-6

LIST OF MAJOR PROCESS EQUIPMENT FOR SILANE PROCESS - CASE B

			Decirima for Sidema 1	recesso cases s		Materials
		Type	Punction	Duty	Size	of Construction
1.	(T1)	M.G. Silicon Storage Hopper	Raw material Storage	2 weeks storage	1.363 x 10 ⁴ gallons	cs
2.	(T2)	Hydrogen Storage Tank	Raw Material Storage	8 hours backup for pipeline failure	9.161 x 10 ⁴ gallons 250 FSIA (spherical)	cs
3.	(T3)	Liquid HCl Storage Tank	Raw Material Storage	2 weeks storage	1.612 x 10 ⁴ gallons 250 PSIA, -50°F (spherical)	Mickel Steel
4.	(T4)	Recycle TET Storage	For TCS REactor Feed	l day storage	9.923 x 10 ⁴ gallons 65 PSIA	cs
45. 4	(T5)	TCS Reactor Off- Gas Flash Tank	Phase Separation		l ft. diameter by 4 long, 65 PSIA, 0°F 65 PSIA	ft. CS
6.	(T6)	TCS/TET Storage	Feed Distillation Column #1	l day hold-up	1.966 x 10 ⁵ 65 PSIA	CS
7.	(T7)	#1 Distillation Column Condensate Accumulator	Reflux feed; column Control	20 minutes hold-up	4.88 x 10 ³ gallons 65 PSIA	CS
8.	(T8)	#2 Distillation Column Condensate Accumulator	Reflux feed; column Control	20 minutes hold-up	746 gallons 65 PSIA	SS
9.	(T9)	#3 Distillation Column Condensate Accumulator	Reflux feed; phase Separation; column control	20 minutes hold-up	194 gallons 5°F. 60 PSIA	SS

10.	(T10)	#4 Distillation Column Condensate Tank	Reflux feed; column control	20 minutes hold-up	18 gallons 50 PSIA, -7°F	SS
11.	(Tll)	Waste Tank	Collect waste for Treatment and disposal	2 week storage	1.378 x 10 ⁴ gallons 65 PSIA	cs
12.	(T12)	Silane Storage	Final Product Storage	2 days storage 8.97 x 10 ³ BTU/Hr	4.349×10^3 gallons -40° F, 250 PSIA	SS
13.	(T13)	Caustic Storage	Raw Material Storage	2 weeks storage	2.304 x 10 ⁴ gallons	<i>S</i> S
14. #	(H1)	TCS Reactor Recycle Gas Heater	Heat Recycle gas and Hydrogen to 550°C	2.342 x 10 ⁶ BTU/hr	752 ft ² 65 PSIA	cs
⁵ 15.	(H2)	HCl Vaporizer	Heat Reactant to 550°C	1.587 x 10 ⁵ BTU/hr	34 Ft ² 65 PSIA	cs
16.	(H3)	TET Vaporizer	Heat Reactant to 550°C	8.755 x 10 ⁶ BTU/hr	2381 ft ² 65 PSIA	CS
17.	(H4)	TCS Teactor Re- cycle Condenser	Phase separation; Recycle hydrogen	1.094 x 10 ⁷ BTU/hr	1882 ft ² 65 PSIA	CS/SS
18.	(H5)	#1 Distillation Column Preheater	Preheat distillation feed to bubble point	2.044 x 10 ⁶ BTU/hr	164 ft ² 250 PSIA	CS
19.	(H6)	#1 Distilllation Column Condenser	Provide Reflux to Column	1.296 x 10 ⁷ BTU/hr	3189 ft ² 65 PSIA	CS
20.	(H7)	#1 Distillation Column Reboiler	Provide vapor to Column	2.382 x 10 ⁷ BTU/hr	2818 ft ² 250 PSIA	cs

CASE B

TABLE A1.2-6 (Continued)

21.	(H8)	#2 Distillation Column Condenser	Provide Reflux to column	1.96 x 10 ⁶ BTU/hr	956 ft ² 65 PSIA	CS/SS
22.	(H9)	#2 Distillation Column Reboiler	Provide Vapor to column	2.693 x 10 ⁷ BTU/hr	2514 ft ² 250 PSIA	C\$
23.	(H10)	#2 Redistribution Reactor Feed Vaporizer	Vaporize Reactants for Reactor	8.81 x 10 ⁵ BTU/hr	78 ft ² 250 PSIA	CS/SS
24.	(H11)	#3 Distillation Column Condenser	Provide Column Reflux (Partial Condenser)	7.312 x 10 ⁵ BW/hr	593 ft ² 60 PSIA	CS/SS
4 25. 6	(H12)	#3 Distillation Column Reboiler	Provide Vapor to Column	9.64 x 10 ⁵ BTU/hr	84 ft ² 250 PSIA	CS/SS
26.	(H13)	Silane Condenser	Condense Final Pro- duct for storage	4.9 x 10 ⁴ BTU/hr	53 ft ² 250 PSIA	CS/SS
27.	(H14)	#4 Distillation Column Condenser	Provide Reflux	8.71 x 10 ⁴ BTU/hr	84 ft ² 50 PSIA	CS/SS
28.	(H15)	#4 Distillation Column Reboiler	Provide Vapor to Column	1.2 x 10 ⁵ BTU/hr	13 ft ² 250 PSIA	CS/SS
29.	(H16)	Absorber Pre- cooler	Cool TET for absorp- tion column	1.35 x 10 ⁵ BTU/hr	35 ft ² 60 PSIA	CS/SS
30.	(H17)	Nitrogen Heater	Heat Nitrogen to re- generate Charcoal Adsorbers	2.52 x 10 ⁴ BTU/hr	14.1 ft ²	cs
31.	(P1)	TCS Reactor Off Gas Recycle Com- pressor	Circulate Recycle Gas to Reactor	1.36 x 10 ³ SCFM	26.5 Horsepower 75 PSIA Discharge	CS*

CASE B

32.	(P2)	#1 Distillation Column Feed Pump	Feed Column	136.5 gpm	106 PSI; 14.5 BHP	cs*
33.	(P3)	#1 Distillation Column Overheads Pump	Provide Reflux and remove overhead product	244 gpm	92.3 PSI; 22.5 BHP	CS*
34.	(P4)	#1 Distillation Column Bottoms Pump	Remove Bottoms Product to TET storage tank	69 дъя	106 PSI, 7.3 BHP	CS*
35.	(P5)	Process Water Feed Pump	Feed Process Water to Waste Treatment	48.6 gpm	82.5 PSI; 4 BHP	CS*
.68	(P6)	Caustic Feed	Feed Raw Material	l gpm	118 PSI; 1/4 BHP	SS
4 ₁₇ 37.	(P7)	#2 Distillation Column Overheads Pump	Provide Reflux and Remove Overhead Product	37.3 дря	92.3 PSI; 3.4 BPH	SS
38.	(P8)	#2 Distillation Column Bottoms Pump	Remove Bottoms Product to TCS/TET storage tank	66.7 gpma	106.3 PSI; 7.1 BPH	SS
39.	(P9)	#3 Distillation Column Overhead Pump	Provide Reflux; Remove Overhead Product	9.7 gpm	87.3 PSI; 1 BHP	S S
40.	(P10)	#3 Distillation Column Bottoms Pump	Remove Bottoms Product to TCS/TET Tank	5.2 дря	106.3 PSI; 1/2 BHP	SS

^{*} Includes incremental higher cost for special purity requirements.

4	1. (P)	(1) #4 Distillation Column: Overhead Pump	,	lgpm	77.3 PSI; 1/4 BHP	SS
4	2. {P	(2) #4 Distillation Column Bottoms Pump	Remove Bottoms Product to Absorber Feed Tank	ī dipma	91.3 PSI; 1/4 BEP	SS
1	7. (P)	3) Silane Product Compressor	Liquefy Silane for Storage	66 SCPM	250 PSIA Discharge 6.5 HP	SS
:	4. 'P	.4) Waste Feed Pump	Distillation Wastes to Waste Treatment	l ggama	76.3 PSI; 1/4 BEP	cs
<u>-</u> 4	5. (P	5) TCS Peactor Ree Pump	ed Feed TET to Reactor	69 dbar	92.3 PSI; 6.4 BEP	CS*
4	6. P	.6) Waste Collection Pump	on Distillation Wastes to Waste Tank	l gipta	87.3 PSI; 1/4 BEP	cs
-	7. (8)) #1 Distillation Column	Separate TET from TCS	94,220 lb/hr of feed	7.56 ft. diameter 100 ft. tall, 50 tr /s	CS
4	e. (c:	!) #2 Distillation Column	Separate TCS from DCS	48,321 lb/hr of feed	10.6 ft. Diameter 136 ft. tall, 68 trays	cs
4	9. /#:	3) #3 Distillation Column	Separate Silane from other Chlorosilanes	7344 lb/hr of feed	2.01 ft. Diameter 29 ft. tall, 29 trays	55
Š:	0. (C	i) #4 Distillation Column	Strip TET for use in absorber	1007.7 lb/hr of feed	1.04 ft. Diameter 28.5 ft. tall, 38 trays	SS
5	1. (cs) Silame Absorber	Absorb Chlorosilane from Silane	819.3 lb/hr of vapor feed	0.823 ft. Diameter 12 ft. tall, 16 trays	SS

^{*} Includes incremental higher cost for special purity requirements.

TABLE A1.2-6 (Continued)

	52.	(C6)	Charcoal Adsorber	Activated Carbon Adsorb- tion of Silane to remove Trace Chlorosilane	-	l ft. Diameter 7 ft. tall (2), 623 lbs of carbon	SS
	53.	(R1)	TCS Fluidized Bed Reactor	Produces TCS from TET,M.G. Silicon, and H ₂		6.26 ft. in Diameter 26.5 ft. tall, 481 tubes 1", 16' long	5\$
	54.	(R2)	#1 Redistribu- tion Reactor (2)	Redistribute TCS to DCS		2' Diameter by 15 ft. tall 1042 lbs catalyst	SS
	55.	(R3)	#2 Redistribu- tion Teactor (2)	Redistribute L to Silane		2.34' Diameter by 35 ft. tall 1667.2 lbs catalyst	SS
419	56.	(Al)	Fines Separator	Remove Silicon Fines carried over with TCS Reactor Off-gas		Standard design 30" Diameter	SS
	57.	(A2)	Waste Treatment	Discharge innorwous effluent		l column for adsorption ÷ 1 heat exchanger to vapor- ize feed	SS
	58.	(A3)	Hydrogen Flare	Dispose of Hydrogen from Waste Treatment		30 ft. stack 6" Diameter	CS

TABLE Al.2-7

PRODUCTION LABOR REQUIREMENTS FOR SILANE PROCESS - CASE B

	Unit Operation	туре	Skilled L Per Day	abor, Man Hours Fer lb. Silane
1.	TCB Production	В	65	,0085
2.	Hydrogen Recycle	c	18	.0023
3.	Raw Material Vaporization	C	50	.0065
4.	TCS Condensation	С	50	.0065
5.	TCS/TET Separation	С	62	.0081
6.	#1 Redistribution Reactor	C	49	.0064
7.	DCS/TCS Separation	C	52	.0068
8.	#2 Redistribution Reactor	C	32	.0042
9.	Silane Distillation	C	32	.0042
10.	Silane Absorption	С	28	.0036
11.	Silane Purification (adsorption)	n	36	.0047
12.	Silane compression	B	23	.003
13.	Silane Condensation	В	23	.003
14.	Materials Handling	A	48*	.0063*
15.	Waste Treatment	В	60	.0078
16.	Silicon Fines Separation	A	15	.002
	TOIA!		 95	. 0776

NOTEs:

^{1.} A Batch Process of Miltiple Small Units

B Average Process

C Automated Process

^{2.} Man hours day dood from Figure 4-6, Poter, and Fimmerhaus (7).

A1.3 Silane Process - Case C

Initial results for the Silane Process (Cases A and B) were marginal and indicated process revisions were warranted.

Based on these initial findings, Union Carbide engineering, research development personnel revised their flowsheet for a more optimum arrangement of major process equipment, raw material requirements and operating conditions. A joint meeting with Union Carbide and Lamar was conducted in late January (1978) for initial review of the revised flowsheet and potential lower plant capital investment and lower product cost for silane production.

In the revised silane process, the silicon tetrachloride is hydrogenated in a fluidized bed of silicon which is catalyzed by copper. The hydrogenation reaction is conducted at a higher pressure than originally proposed to increase the yield of desireable trichlorosilane. The gas leaving the fluidized bed reactor is cooled and condensed to recover the liquid chlorosilanes. The hydrogen is recycled.

The condensed liquid chlorosilanes are separated by distillation. The inerts (dissolved gases) are removed in the initial distillation column. The remaining distillation columns separate the liquid chlorosilanes into primarily solicon tetrachloride, trichlorosilane, dichlorosilane and silane. The silicon tetrachloride is recycled back to the hydrogenation reactor. The trichlorosilane and dichlorosilane are sent to the redistribution reactors for rearrangement of chlorine/hydrogen bonds to silicon. The final redistribution reactor product is sent to the silane distillation column. The silane is removed from this distillation and sent to silicon production.

Chemical engineering analysis results for the Silane Process - Case C (Revised Process) are given in Section 3.3- UCC Silane Process for Silicon (Union Carbide Corporation).

A2. ADDITIONAL ECONOMIC ANALYSIS

A2.1 Silane Process - Case A

The economic analysis activity for the Silane Process
-Case A (Regular Process Storage) involves a cost analysis
to produce silane for silicon. Primary results issuing from
the economic analysis include plant capital investment and
product cost which are useful in identification of those processes showing promise for meeting project cost goals.

The cost analysis results for producing silane by the Silane Process - Case A are presented in Table A2.1-1 including costs for raw materials, labor, utilities and other items composing the product cost (total cost of producing silicon). The tabulation summarizes all of these items to give a total product cost without profit of \$5.55 (1975 dollars) and \$7.77 (1980 dollars) per 1b of silane. This product cost without profit includes direct manufacturing cost, indirect manufacturing cost, plant overhead and general expenses. These results, when expressed in terms of silicon contained in the silane, correspond to \$13.94 (1975 dollars) and \$19.53 (1980 dollars) per kg of silicon.

This cost results for the Silane Process - Case A indicate that this new technology for producing silane for silicon is marginal. Revisions are warranted for meeting the cost goal of \$14 per kg of silicon material (1980 dollars) for solar cells.

The detailed results for the economic analysis are presented in a tabular format to make it easier to locate cost items of specific interest. The guide for the tabular format is given below:

Preliminary Economic Analysis Activities Table	A2.1-2
Process Design Inputs	A2.1-3
Base Case Conditions Table	A2.1-4
Raw Material Cost	A2.1-5
Utility CostTable	A2.1-6
Major Process Equipment Cost	
Production Labor Cost	A2.1-8
Plant Investment	A2.1-9
Total Product CostTable	A2,1-10

TABLE A2.1-1

BETIMATION OF PRODUCT COST FOR SILANS PROCESS - CASE A

		Cost \$/lb of Silane (1975 dollars)	Cost \$/lb of Bilane (1980 dollars)
1.	Direct Manufacturing Cost (Direct Costs) Raw Materials Direct Operating Labor Utilities Supervision and Clerical Maintenance and Repairs Operating Supplies Laboratory Charge	. 3.34	4.68
2.	Indirect Manufactuing Cost (Fixed Cost) Depreciation Local Taxes Insurance	. 0.89	1.25
3.	Plant Overhead	. 0.60	.84
4.	General Expenses	0.72	1.01
5.	Product Cost Without Profit	5.55	7.77

Table A2.1-2

ECONOMIC ANALYSES:

THELIMINARY ECONOMIC ANALYSIS ACTIVITIES FOR SILAME PROCESS

CASE A

		Frel. Process Economic Activity	Status	Prel. Process Economic Activity	States
	1.	Process Design Inputs	6	6. Production Labor Obsts	•
		1. Raw Material Requirements	t	1. Base Cost Per Man Hour	•
		2. Utility Requirements	•	2. Cost/1b Silame Per Area	•
		3. Equipment hist	•	 Total Cost/lb Silame 	•
		4. Labor Requirements	•		
				 Estimation of Plant Investment 	•
	2.	Specify Base Case Conditions		 Battery Limits Direct Costs 	•
		1. Base Year for Costs	•	2. Other Direct Costs	•
		2. Appropriate Indice: for Costs	•	3. Indirect Costs	•
		3. Additional	•	4. Contingency	•
				5. Total Plant Investment	•
41.	3.	Raw Material Costs	•	(Fixed Capital)	
V V		1. Base Cost, Lb. of Material	•	-	
		2. Material Cost/lb of Silane		8. Estimation of Total Product Cost	•
		3. Total Cost/lb of Silane	•	1. Direct Manufacturing Cost	•
				2. Indirect Manufacturing Cost	•
	4.	Utility Costs	•	3. Plant Overhead	•
		1. Base Cost for Each Utility	•	4. By-Product Credit	•
		2. Utility Cost/lb of Silane	•	5. General Expenses	•
		. Total Cosc/lb of Silane	e	6. Total Cost of Product	•
	5.	Major Process Equipment Costs	•		
		 Individual Equipment Cost 	•		
		2. Cost Index Adjustment	•		
				0 Plan	
				<pre>9 In Progress</pre>	
				• Complete	

TABLE A2.1-3

PROCESS DESIGN INPUTS FOR SILAME PROCESS - CASE A

- 1. Raw Material Requirements
 - -M.G. Silicon, anhydrous HCl, caustic, hydrogan.
 - -see table for "Raw Material Cost"
- 2. Utility
 - -electrical, steam, cooling water, etc.
 - -see table for "Utility Cost"
- 3. Equipment List
 - -76 pieces of major process equipment
 - -process vessels, heat exchangers, reactor, etc.
 - -see table for "Major Process Equipment Cost"
- 4. Labor Requirements
 - -production labor for purification, vaporization, product handling, etc.
 - -see table for "Production Labor Cost"

CASD A

TABLE A2.1-4

BASE CASE CONDITIONS FOR SILANE PROCESS-CASE A

- 1. Capital Equipment
 - -January 1975 Cost Index for Capital Equipment Cost
 - -January 1974 Cost Index Value = 430
- 2. Utilities
 - Electrical, Steam, Cooling Water, Nitrogen
 - -January 1975 Cost Index (U.s. Dept. Labor)
 - -Values determined by literature search and summarized in cost standardization work
- 3. Paw Material Cost
 - -Chemical Marketing Reporter
 - -January 1975 Value
 - -Other Sources
- 4. Labor Cost
 - -Average for Chemical Petroleum, Comb and Allied Industries (1975)
 - -SFilled \$6.90/hr
 - -Semiski'led \$4.90/hr
- 5. Update to 1980
 - historically cite 1975 dollars (LSA project)
 - 1911. decision to change to 1980 dollars (JPL, 6/22/79)
 - -reports to reflect both 1975 and 1980 dollars (JPL, 6/22/79)
 - -inflation factor of 1.4 to be used (JPL, 6/22/19)

CASE A

TABLE A2.1-5

RAW MATERIAL COST FOR SILANE PROCESS - CASE A

Raw Material	Requirement 1b/1b of Silane	\$/lb of Material	Cost \$/lb of Silane
1. HCl	1.239	.10	.12
2. Hydrogen	.362	.96	.35
3. Caustic (50%)	2.448	.0382	.09
4. M.G. Silicon	1.11	.454	.50
			1.06 (1975 dollar.) x 1.4 inflation 1.48 (1980 dollars)

TABLE A2.1-6 UTILITY COST FOR SILANE PROCESS -CASE A

<u> </u>	tility	Requirement/lb Cost of or Silane Utility	Cost \$/lb of Silane
1.	filectricty	.253 Kw-Hr \$.03/KW hr	.0076
2.	Steam	190.34 Ibs 1.25/M Ib	.2379
3	Cooling Water	168,12 gull ma .09/d gal	.0134
4.	Process water	8.22 gallons .35/M gal	.0029
5.	Rofrigerant (23°F)	27 1 RTU 4.7 /NM BTU	.0001
б.	Refrigerant (5°F)	79.1 RM (.40, EM PMU	.0005
7.	Refrigerant (-7°F)	26.4 BTU 7.50/MM BTU	.0002
8.	Refrigerant (-20°F)	2.3 M BTU 8.70/MM BTU	.0200
•	Refrigerant (30°F)	30 8 M BTH O GO, MM BTH	.2957
19.	Refrigerant (-40°F)	269 BTH 10.50/MM PTH	· 0039
11.	Refrigerant (-50°F)	3.5 M BRU 11.47/MM RT.	.0400
12.	High Temperature Heat	3.324 x 10 1 BTU 3.0 MM BTU	.099
13.	Mkuh inge Pluid Ni drogan	5.54 SCF .50, M SCF	.0028
			.724 (1975 dollars) x 1.4 inflation

1.01 (1980 dollars)

TABLE A2.1-7

PURCHASED COST OF MAJOR PROCESS EQUIPMENT FOR SILANE PROCESS - CASE A

1.	Equip (T1)	M.G. Silicon Storage Hopper	Purchased Cost,\$1000 12.05
2.	(T2)	Hydrogen Storage Tank	179.2
3.	(T3)	Liquid HCl Storage Tank	95.27
4.	(T4)	Recycle TET Storage	214,4
5.	(T5)	TCS Reactor Off-Gas Flash Tank	0.71
6.	(T6)	TCS/TET Storage	214.4
7.	(T7)	#1 Distillation Column Condensate Accumlator	8.51
8.	(8T)	#1 Rediscribution Reactor Feed Tank	241.99
9.	(T9)	#1 Redistribution Reactor Product Tank	245.0
10.	(T10)	#2 Distillation Column Condensate Accumulator	7.37
11.	(T11)	#2 Redistribution Reactor Feed Tank	76.03
12.	(T12)	#2 Redistribution Reactor Product Tank	221.17
13.	(T13)	#3 Distillation Column Condensate Accumulator	2.76
14.	(T14)	#3 Distillation Column Condensate Tank	147.44
15.	(T15)	#4 Distillation Column Feed Tank	53.45
16.	(T16)	#4 Distillation Column Condensate Accumulator	2.76
17.	(T17)	#4 Distillation Column Condensate Tank	34.1
18.	(T18)	Waste Tank	17.01
19.	(T19)	Absorber Feed Tank	16.59
20.	(T20)	Silane Storage	255.9
21.	(T21)	Caustic Storage	92.15
22.	(H1)	TCS Reactor Recycle Gas Heater	8.12
23.	(H2)	HCl Vaporizer	1.15
24.	(H3)	TET Vaporizer	18.48
25.	(H4)	TCS Reactor Recycle Condenser	38.98

TABLE A2.1-7 (continued)

26.	(H5)	#1 Distillation Column Proheater	3.24
27.	(H6)	#1 Distillation Column Condenser	22.4
28.	(H7)	#1 Divillation Column Reboiler	23.7
29.	(H8)	#2 Distillation Column Condenser	21.08
30.	(H9)	#2 Distillation Column Reboiler	21.16
31.	(H10)	#2 Redistribution Reactor Food Vaporiser	3.67
32.	(H11)	#2 Redistribution Reactor Product condenser	8.62
33.	(H12)	#3 Distillation Column Preheater	2.86
34.	(H13)	#3 Distillation Column Condenser	14.95
35.	(H14)	#3 Distillation Column Reboiler	3.88
36.	(H15)	Silane Condenser	2,29
37.	(H16)	#4 Distillation Column Condenser	3.48
38.	(H17)	#4 Distillation Column Reboiler	1.33
39.	(H18)	Absorber Pre-cooler	1.79
40.	(H19)	Nitrogen Heater	.92
41.	(Pl)	TCS Reactor Off-gas Recycle Compressor	35.1
42.	(P2)	#1 Distillation Column Feed Pump	5.03
43.	(P3)	#1 Distillation Column Overheads Pump	6.04
44.	(P4)	#1 Distillation Colum Bottoms Pump	3.59
45.	(P5)	Process Water Feed Pump	2.87
46.	(P6)	Caustic Feed Pump	1.25
47.	(P7)	#1 Redistribution Reactor Feed Pump	4.02
48.	(P8)	#2 Distillation Column Feed Pump	3.59
49.	(F9)	#2 Distillation Column Overheads Pump	2.57
50.	(P10)	#2 Distillation Column Bottoms Pump	3.59
51.	(P11)	#2 Redistribution Reactor Feed Pump	2.09
52.	(P12)	#3 Distillation Column Feed Pump	1.77

TABLE A2.1-7 (continued)

53.	(P13)	#3 Distillation Column Overheads Pump	1.77
54.	(P14)	#3 Distillation Column Bottoms Pump	1.47
55.	(PI5)	#4 Distillation Column Feed Pump	1.23
56,	(P16)	#4 Distillation Column Overheads Pump	1.23
57.	(P17)	#4 Distillation Column Bottoms Pump	1.23
58.	(P18)	#4 Distillation Condensate Recycle Pump	1.23
59.	(r19)	Silane Product Compressor	17.55
60.	(P20)	Waste Feed Pump	.62
61.	(P21)	TCS Reactor Feed Pump	3.31
62,	(P22)	#3 Dis lation Condensate Recycle Pump	1.47
63.	(P23)	Waste Collection Pump	.62
64.	(P24)	Absorber Feed Pump	1.23
65.	(C1)	#1 Distillation Column	100.66
66,	(C2)	#2 Distillation Column	214.08
67.	(C3)	#3 Distillation Column	40.19
68.	(C4)	#4 Distillation Column	21.14
69.	(C5)	Silane Absorber	15.06
70,	(ce)	Charcoal Adsorber	18.0
71,	(R1)	TCS Fluidized Bed Reactor	155.06
72.	(R2)	#1 Redistribution Reactor	13.26
73.	(R3)	#2 Redistribution Reactor	33.14
74.	(A1)	Fines Separator	2.0
75.	(A2)	Waste Treatment	18.72
76.	(F.A)	Hydrogen Flare	0.10
		TOTAL PURCHASED EQUIPMENT COST	\$3079.31 (1975 dollars) x 1.4 inflation 4,311.03 (1980 dollars)

CASE A

TABLE A2.1-8

PRODUCTION LABOR COST FOR SILANE PROCEES - CASE A

	Unit Operation	Skilled Labor Man-Hrs/1b Silane	Cost \$/1b of Silane
1.	TCS Production	.0085	.05865
2.	Hydrogen Recycle	.0023	.01587
3.	Raw Material Vaporization	.0065	. 04485
4.	TCS Condensation	.0065	.04485
5.	TCS/TET Separation	.0081	.05589
6.	#1 Redistribution Reactor	.0064	.04416
7.	DCS/TCS Separation	.0068	.04692
8.	#2 Redistribution Reactor	.0042	.02898
9.	Silane Distillation	.0042	.02898
10.	Silane Absorption	.0036	.02484
11.	Silane Purification (Adsorption)	.0047	.03243
12.	Silane Compression	.003	.0207
13.	Silane Condensation	.003	.0207
14.	Materials Handling	.063*	.03087
15.	Waste Treatment	.0078	.05382
16.	Silicon Fines Separation	,002	.0138 \$.5663 (1975 dollars) x 1.4 inflation .7928 (1980 dollars)

NOTES

Based on labor costs of \$6.90 skilled, \$4.90 semiskilled.

^{*} Semiskilled Labor

TABLE A2.1-9

BETIMATION OF PLANT INVESTMENT FOR SILANE PROCESS - CASE A

		Investment \$1000	
1.	DIRECT PLANT INVESTMENT COSTS		
	1. Major Process Equipment Cost	\$3079.31	
	2. Installation of Major Process Equipment	1324.10	
	3. Process Piping, Installed	2278.69	
	4. Instrumentation, Installed	585.07	
	5. Electrical, Installed	307.93	
	6. Process Buildings, Installed	307.93	
la.	SUBTOTAL FOR DIRECT PLANT INVESTMENT COSTS	7883.03	
	(PRIMARILY BATTERY LIMIT PACILITIES)		
2.	OTHER DIRECT PLANT INVESTMENT COSTS		
	1. Utilities, Installed	1478.07	
	2. General Services, Site Development,	369.52	
	Fire Protection, etc.		
	3. General Buildings, Offices, Shops, etc.	431.10	
	4. Receiving, Shipping Facilities	646.66	
2a.	SUBTOTAL FOR OTHER DIRECT PLANT INVESTMENT COUTS (PRIMARILY OFFSITE FACILITIES OUTSIDE BATTERY LIMITS)	2925.35	
3.	TOTAL DIRECT PLANT INVESTMENT COST, la + 2a	10808.38	
4.	INDIRECT PLANT INVESTMENT COSTS	3.605.60	
	1. Engineering, Overhead, etc.	1693.62	
	2. Normal Cont. for Floods, Strikes, etc.	2186.31	
4a.	TOTAL INDIRECT PLANT INVESTMENT COST	3879.91	
5,	TOTAL DIRECT AND INDIRECT PLANT INVESTMENT COST, 3 + 4a	14688.31	
6.	OVERALL CONTINGENCY	4406.49	
7.	FIXED CAPITAL INVESTMENT FOR PLANT, 5 + 6	10004 00	/1076 3:33
• •	TARRE OF THE TANDIDAT DATE THAT! O T O		(1975 dollars)
			inflation
		26/32.72	(1980 dollars)

TABLE A2.1-10

ESTIMATION OF TOTAL PRODUCT COST FOR SILANE PROCESS- CASE A

		\$/1b of Silane
1.	Direct Manfacturing Cost (Direct Charges)	
	1. Raw Materials- from prel. design	1.06
	2. Direct Operating Labor- from prel.	.5663
	3. Utilities-from prel. design	.724
	4. Supervision and Clerical,	.085
	5. Maintenance and Repairs,	.682
	6. Operating Supplies,	.136
	7. Laboratory Charge,	.085
	8. Patents and Royalties, costs	का क्ष्म का <i>क</i>
2.	Indirect Manufacturing Cost (Fixed Charges)	
	1. Depreciation	.682
	2. Local Taxes	.136
	3. Insurance	.068
	4. Interest	ps par lest 400
3.	Plant Overhead	.595
4.	By-Product Credit- from prel. design	
4a.	Total Manufacturing Cost, 1 + 2 + 3 + 4	4.819
5.	General Expenses	.289
	1. Administration,	.289
	2. Distribution and Sales,	. 209
	3. Research and Development, cost	,145
6.	Total Cost of Product, 4a + 5	5.55 (1975 dollars) x 1.4 inflation 7.77 (1980 dollars)

A2.2 Silane Process - Case B

The economic analysis activity for the Silane Process
-Case B (Minimum Process Storage) involves a cost analysis
to produce silane for silicon. Primary results issuing from
the economic analysis include plant capital investment and
product cost which are useful in identification of those pro-esses showing promise for meeting project cost goals.

The cost analysis results for producing silane by the Silane Process - Case B are presented in Table A2.2-1 including costs for raw materials, labor, utilities and other items composing the product cost (total cost of producing silicon). The tabulation summarizes all of these items to give a total product cost without profit of \$4.58 (1975 dollars) and \$6.41 (1980 dollars) per 1b of silane. These results, when expressed in terms of silicon contained in the silane, correspond to \$11.53 (1975 dollars) and \$16.12 (1980 dollars) per kg of silicon.

These cost results for the Silane Process-Case B indicate that this new technology for producing silane for silicon is marginal. Revisions are warranted for meeting the cost goal of \$14 per kg of silicon material (1980 dollars) for solar cells.

The detailed results for the economic analysis are presented in a tabular format to make it easier to locate cost items of specific interest. The guide for the tabular format is given below:

Preliminary Economic Analysis Activities Table	A2,2-2
Process Design Inputs	A2.2-3
Base Case Conditions	A2, 2-4
Raw Material Cost	A2.2-5
Utility CostTable	A2.2-6
Major Process Equipment	A2.2-7
Production Labor Cost	A2.2-8
Plant Investment	A2,2-9
Total Product Cost	A2.2-10

TABLE A2.2-1

ESTIMATION OF PRODUCT COST FOR SILANE PROCESS - CASE B

		Cost \$/lb of Silane (1975 dollars)	Cost \$/lb of Silane (1980 dollars)
1.	Direct Manufacturing Cost (Direct Costs) Raw Materials Direct Operating Labor Utilities Supervision and Clerical Maintenance and Repairs Operating Supplies Laboratory Charge	. 2.95	4.13
2.	Indirect Manufactuing Cost (Fixed Cost) Depreciation Local Taxes Insurance	, 0.52	0.73
з.	Plant Overhead	0.51	0.71
4.	General Expenses	0.60	0.84
5.	Product Cost Without Profit	4.58	6.41

TABLE A2.2-2

ECONOMIC ANALYSES:

PRELIMINARY ECONOMIC ANALYSIS ACTIVITES FOR SILANE PROCESS - CASE B (UNION CARBIDE)

		Prel. Process Economic Activity	<u>Status</u>		Prel. Process Economic Activity	States
	ì.	Process Design Inputs	•	6.	Production Labor Costs	•
		1. Faw Material Requirements	9		1. Base Cost Per Man Hour	•
		2. Utility Requirements	\$		Cost/lb Silane Per Area	•
		3. Equipment List	•		Total Cost/lb Silane	•
		4. Labor Requirements	•			
				7.	Estimation of Plant Investment	•
		Specify Base Case Conditions			1. Battery Limits Direct Costs	•
		1. Base Year for Costs	•		2. Other Direct Costs	
		2. Appropriate Indices for Costs	5		3. Indirect Costs	•
		3. Additional	•		4. Contingency	•
					5. Total Plant Investment	•
. <u>4</u> . ω	3.	Paw Material Costs	•		(Fixed Capital)	
37		 Base Cost/Lb. of Material 	•		- ·	
		2. Material Cost/lb of Silane	•	8.	Estimation of Total Product Cost	•
		3. Total Cost/lb of Silane	•		l. Direct Manufacturing Cost	•
					2. Indirect Manufacturing Cost	•
	4.	Utility Costs	•		3. Plant Cwrhead	
		1. Base Cost for Each Utility	•		4. By-Product Credit	•
		2. Utility Cost/lb of Silane	•		5. General Expenses	•
		3. Total Cost/lb of Silane	•		6. Total Cost of Product	•
	5.	Major Process Equipment Costs	•			
		 Individual Equipment Cost 	•			
		Cost Index Adjustment	•			
					0 Plan	
					0 In Progress	
					• Complete	

TABLE A2.2-3

PROCESS DESIGN INPUTS FOR SILANS PROCESS - CASE B

- 1. Raw Material Requirements
 - -M.G. Silicon, anhydrous HCl, caustic, hydrogen.
 - -see table for "Raw Material Cost"
- 2. Utility
 - -electrical, steam, cooling water, etc.
 - -see table for "Utility Cost"
- 3. Equipment List
 - -58 pieces of major process equipment
 - -process vessels, heat exchangers, reactor, etc.
 - -see table for "Major Process Equipment Cost"
- 4. Labor Requirements
 - -production labor for purification, vaporization, product handling, etc.
 - -see table for "Production Labor Cost"

TABLE A2.2-4

BASE CASE CONDITIONS FOR SILANE PROCESS - CASE B

1. Capital Equipment

- -January 1975 Cost Index for Capital Equipment Cost
- -January 1975 Cost Index Value = 430

2. Utilities

- -Electrical, Steam, Cooling Water, Nitrogen
- -January 1975 Cost Index (U.S. Dept. Labor)
- -Values determined by literature search and summarized in cost standardization work

3. Raw Material Cost

- -Chemical Marketing Reporter
- -January 1975 Value
- -Other Sources

4. Labor Cost

- -Average for Chemical Petroleum, Coal and Allied Industries (1975)
- -Skilled \$6.90/hr
- -Semiskilled \$4.90/hr

5. Update to 1980

- -historically cited 1975 dollars (LSA project)
- -DOE decision to change to 1980 dollars (JPL, 6/22/79)
- -reports to reflect both 1975 and 1980 dollars (JPL, 6/22/79)
- -inflation factor of 1.4 to be used (JPL, 6/22/79)

CASE B

TABLE A2.2-5 RAW MATERIAL COST FOR SILAME PROCESS-CASE B

Raw	Matorial	Requirement 1b/1b of Silane	\$/1b of Material	Cost \$/lb of Silane
1.	нст	1.239	.10	.12
2.	Hydrogen	.362	.96	.35
3.	Caustic (50%)	2.448	.0382	.09
4.	M.G. Silicon	1.11	.454	.50
				1.06 (1975 dollars) x 1.4 inflation 1.48 (1980 dollars)

CASE B

TABLE A2.2-6 UTILITY COST FOR STLANE PROCESS -CASE B

Utility		Requirement/lb of Silane	Cost of Utility	Cost \$/lb of Silane	
1.	Electricty	.212 KW-Hr	\$.03/kw hr	.0064	
2.	Steam	186.72 lbs	1.25/M lb	. 2334	
3.	Cooling Water	168.12 g&llons	.08/M gal	.0134	
4.	Process Water	8.22 gallons	.35/M gal	.0029	
5.	Refrigerant (-20°F)	2.3 M BTU	8.70/MM BTU	.0200	
6.	Refrigerant (-30°F)	30.8 M BTU	9.60/MM BTU	.2957	
7.	Refrigerant (-40°F)	25.3 BTU	10.50/MM BTU	.0003	
8.	Refrigerant (-50°F)	517.2 BTU	11.42/MM BTU	.0059	
9.	High Temperature Heat Exchange Fluid	33,24 M BTU	3.0/MM BTU	.0997	
10.	Nitrogen	5.54 SCF	.50/M SCF	.0028	
				.6805 (1975 dollars)	

x 1.4 inflation .9527 (1980 dollars)

TABLE A2.2-7

PURCHASED COST OF MAJOR PROCESS EQUIPMENT FOR SILANE PROCESS - CASE B

	Equip	ment	Purchased Cost, \$1000
1.	(T1)	M.G. Silicon Storage Hopper	12.05
2.	(T2)	Hydrogen Storage Tank	179.2
3.	(T3)	Liquid HCl Storage Tank	95.27
4.	(T4)	Recycle TET Storage	125.55
5.	(T5)	TCS Reactor Off-Gas Flash Tank	0.71
6.	(T6)	TCS/TET Storage	214.4
7.	(T7)	#1 Distillation Column Condensate Accumulator	8.51
8.	(T8)	#2 Distillation Column Condensate Accumulator	7.37
9.	(T9)	#3 Distillation Column Condensate Accumulator	2.76
10.	(T10)	#4 Distillation Column Condensate Accumulator	2.76
11.	(T11)	Waste Tank	17.01
12.	(T12)	Silane Storage	82.09
13.	(T13)	Caustic Storage	92.15
14.	(H1)	TCS Reactor Recycle Gas Heater	8.12
15.	(H2)	HCl Vaporizer	1.15
16.	(H3)	TET Vaporizer	18.48
17.	(H4)	TCS Reactor Recycle Condenser	38.98
18.	(H5)	#1 Distillation Column Preheater	3.24
19.	(H6)	#1 Distillation Column Condenser	22.4
20.	(H7)	#1 Distillation Column Reboiler	23.7
21.	(H8)	#2 Distillation Column Condenser	21.08

TABLE A2.2-7 (Continued)

22.	(<i>P</i> 9)	#2 Distilation Column Reboiler	21.16
23.	(H1O)	#2 Redistribulton Reactor Feed Vaporizer	2.67
24.	(H11)	#3 Distillation Column Condenser	14.95
25.	(H12)	#3 Distillation Column Reboiler	3.88
26.	(H13)	Silame Condenser	2.29
27.	(H14)	#4 Distillation Column Condenser	3.48
28.	(B15)	#4 Distillation Column Reboiler	1.33
29.	(H16)	Absorber Pre-cooler	1.79
30.	(H17)	Nitrogen Heater	.92
31.	(P1)	TCS Reactor Off-gas Recycle Compressor	35.1
32.	(P2)	#1 Distillation Column Feed Pump	5.03
33.	(P3)	#1 Distillation Column Overheads Pump	6.04
34.	(P4)	#1 Distillation Column Bottoms Pump	3.59
35.	(P5)	Process Water Feed Pump	2.87
36.	(P6)	Caustic Feed Pump	1.25
37.	(P7)	#2 Distillation Column Overheads Pump	2.57
38.	(84)	#2 Distillation Column Bottoms Pump	3.59
39.	(P9)	#3 Distillation column Overheads Pump	1.77
40.	(P10)	#3 Distillation Column Bottoms Pump	1.47
41.	(P11)	#4 Distillation Column Overheads Pump	1.23
42.	(1,15)	#4 Distillation Column Bottoms Pump	1.23
43.	(513)	Silane Product Compressor	17.55
44.	(P14)	Waste Feed Pump	.62

TABLE A2.2-7 (Continued)

45.	(P15)	TCS Reactor Feed Pump	3.31
46.	(P16)	Waste Collection Pump	.62
47.	(C1)	#1 Distillation Column	100.66
48.	(C2)	#2 Distillation Column	214.08
49.	(C3)	#3 Distillation Column	40.19
50.	(C4)	#4 Distillation Column	21.14
51.	(C5)	Silane Absorber	15.06
52.	(C6)	Charcoal Adsorber	18.0
53.	(R1)	TCS Fluidized Bed Reactor	155.06
54.	(R2)	#1 Redistribuiton Reactor	26.52
55.	(R3)	#2 Redistribution Reactor	66.28
56.	(A1)	Fines Separator	2.0
57.	(A2)	Waste Treatment	18.72
58.	(A3)	Hydrogen Flare	0.10

TOTAL PURCHASED EQUIPMENT COST 1796.17 (1975 dollars) $\frac{\times 1.4}{2515}$ (1980 dollars)

TABLE A2.2-8

PRODUCTION LABOR COST FOR SILANE PROCESS - CASE B

	Unit Operation	Skilled Labor Man-Hrs/lb Silane	Cost \$/lb of Silans
1.	TCS Production	.0085	.05865
2.	Hydrogen Recycle	.0023	.01587
3.	Raw Material Vaporization	.0065	.04485
4.	TCS Condensation	.0065	.04485
5.	TCS/TET Separation	.0081	.05589
6.	#1 Redistribution Reactor	.0064	.04416
7.	DCS/TCS Separation	.0068	.04692
8,	#2 Redistribution Reactor	.0042	,02898
9.	Silane Distillation	.0642	.02898
10.	Silane Absorption	.0036	.02484
11.	Silane Purification(Adsorption)	.0047	.03243
12.	Silane Compression	.003	.0207
13.	Silane Condensation	.003	.0207
14.	Materials Handling		.03087
15.	Waste Treatment	.0078	.05382
16.	Silicon Fines Separation	.002	.01.38

TOTAL COST .5663 (1975 dollars)
x 1.4 inflation
.7928 (1980 dollars)

NOTES

Based on labor costs of \$6.90 skilled, \$4.90 semiskilled.

^{*} Semiskilled labor.

TABLE A2.2-9

ESTIMATION OF PLANT INVESTMENT FOR SILANE PROCESS - CASE B

		Investment \$1000
1.	DIRECT PLANT INVISTMENT COSTS 1. Major Process Equipment Cost 2. Installation of Major Process Equipment 3. Process Piping, Installed 4. Instrumentation, Installed 5. Electrical, Installed 6. Process Buildings, Installed	\$ 1796.17 772.35 1329.17 341.27 179.62 179.62
la.	SUBTOTAL FOR DIRECT PLANT INVESTMENT COSTS (PRIMARILY BATTERY LIMIT FACILITIES)	4598.2
2.	OTHER DIRECT PLANT INVESTMENT COSTS 1. Utilities, Installed 2. General Services, Site Development, Fire Protection, etc. 3. General Buildings, Offices, Shops, etc. 4. Receiving, Shipping Facilities	862.16 215.54 251.46 377.20
2 a .	SUBTOTAL FOR OTHER DIRECT PLANT INVESTMENT COSTS (PRIMARILY OFFSITE FACILITIES OUTSIDE BATTERY LIMITS)	1706.36
3.	TOTAL DIRECT PLANT INVESTMENT COST, la + 2a	6304.56
4.	INDIRECT PLANT INVESTMENT COSTS 1. Engineering, Overhead, etc. 2. Normal Cont. for Floods, Strikes, etc.	987.89 1275.28
4a.	TOTAL INDIRECT PLANT INVESTMENT COST	2263.17
5.	TOTAL DIPECT AND INDIRECT PLANT INVESTMENT COST, 3 + 4a	8567.73
6.	OVERALL CONTINGENCY	2570.32
7.	FIXED CAPITAL INVESTMENT FOR PLANT, 5 + 6	11138.05 (1975 dollars) x 1.4 inflation 15593 (1980 dollars)

TABLE A2.2-10

ESTIMATION OF TOTAL PRODUCT COST FOR SILANE PROCESS -CASE B

		\$/1b of Silans
1.	Direct Manfacturing Cost (Direct Charges)	
	1. Raw Materials- from prel. design	1.06
	 Direct Operating Labor- from prel. design 	. 5663
	3. Utilities-from prel. design	.6805
	4. Supervision and Clerical,	.0849
	5. Maintenance and Repairs,	. 3976
	6. Operating Supplies,	.0795
	7. Laboratory Charge,	.0849
	8. Patents and Royalties, costs	ope trid dark gam
2.	Indirect Manufacturing Cost (Fixed Charges)	
	1. Depreciation	.3976
	2. Local Taxes	.0795
	3. Insurance	.0398
	4. Interest	one are the face one
3.	Plant Overnead	.51
4,	By-Product Credit- from prel. design	fra ma
4a.	Total Manufacturing Cost, 1 + 2 + 3 + 4	3.9806
5.	General Expenses	955
	1. Administration, cost	.2388
	2. Distribution and Sales,	.2388
	3. Research and Development, cost	.1194
6.	Total Cost of Product, 4a + 5	
		4.58 (1975 dollars)
		\times 1.4 inflation
		6.41 (1980 dollars)

A2.3 Silane Process - Case C

Initial cost analysis results for the Silane Process (Cases A and B) were marginal and indicated process revisions were warranted for meeting the project cost goals.

Process revisions were accomplished with favorable cost benefits over the orginal scheme.

The revised process included operation of the silicon tetrachloride reaction at higher pressure for increased trichlorosilane yield should lower recycle requirements. Lower recycle requirements will lower capital equipment and labor costs. The distillation train as now proposed will operate at several hundred pounds pressure compared to original lower pressure. This higher pressure permits use of cooling water in the condensers and does not require expensive low temperature refrigeration as originally proposed. This will provide lower operating (utilities) cost in 3 of the 4 distillation columns. The higher pressure also permits use of smaller diameter columns (vapor loading, density proportices) to pressure). The elimination of hydrogen chloride reduces starting material costs. Also, the use of hydrogen from the silane pyrolysis provides additional lower feed material costs.

The revised silane process provided the following cost benefits:

lower capital costs

- -lower raw material costs
- -lower operating labor costs

Economic analysis results for the Silane Process - Case C (Revised Process) are given in Section 4.3-UCC Silane Process for Silicon (Union Carbide Corporation).